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(54) **MODULAR ENDOGRAFT DEVICES AND ASSOCIATED SYSTEMS AND METHODS**

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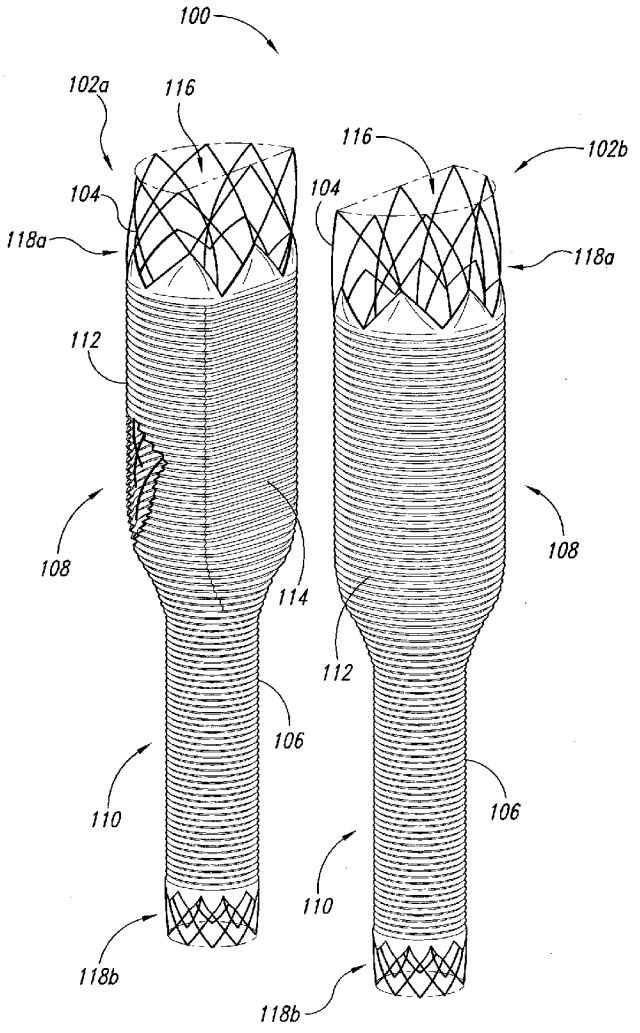
A61F 2/82 (2006.01)

(52) **U.S. Cl.** **623/1.15**

(57) **ABSTRACT**

Modular endograft devices and associated systems and methods are disclosed herein. In several embodiments, an endograft system can include a first endograft device and a second endograft device that each include an integrated frame, a cover and a lumen within the cover. Each endograft device further includes a superior portion and an inferior portion. The superior portion can have a convexly curved outer wall and a septal wall. The first and second endograft devices can be configured to extend into a low-profile configuration with a first cross-sectional dimension and a first length and self-expand into an expanded configuration with a second cross-sectional dimension greater than the first cross-sectional dimension and a second length less than the first length. In the expanded configuration, the septal walls can press against each other and form a septum between the lumens of the first and second endograft devices.



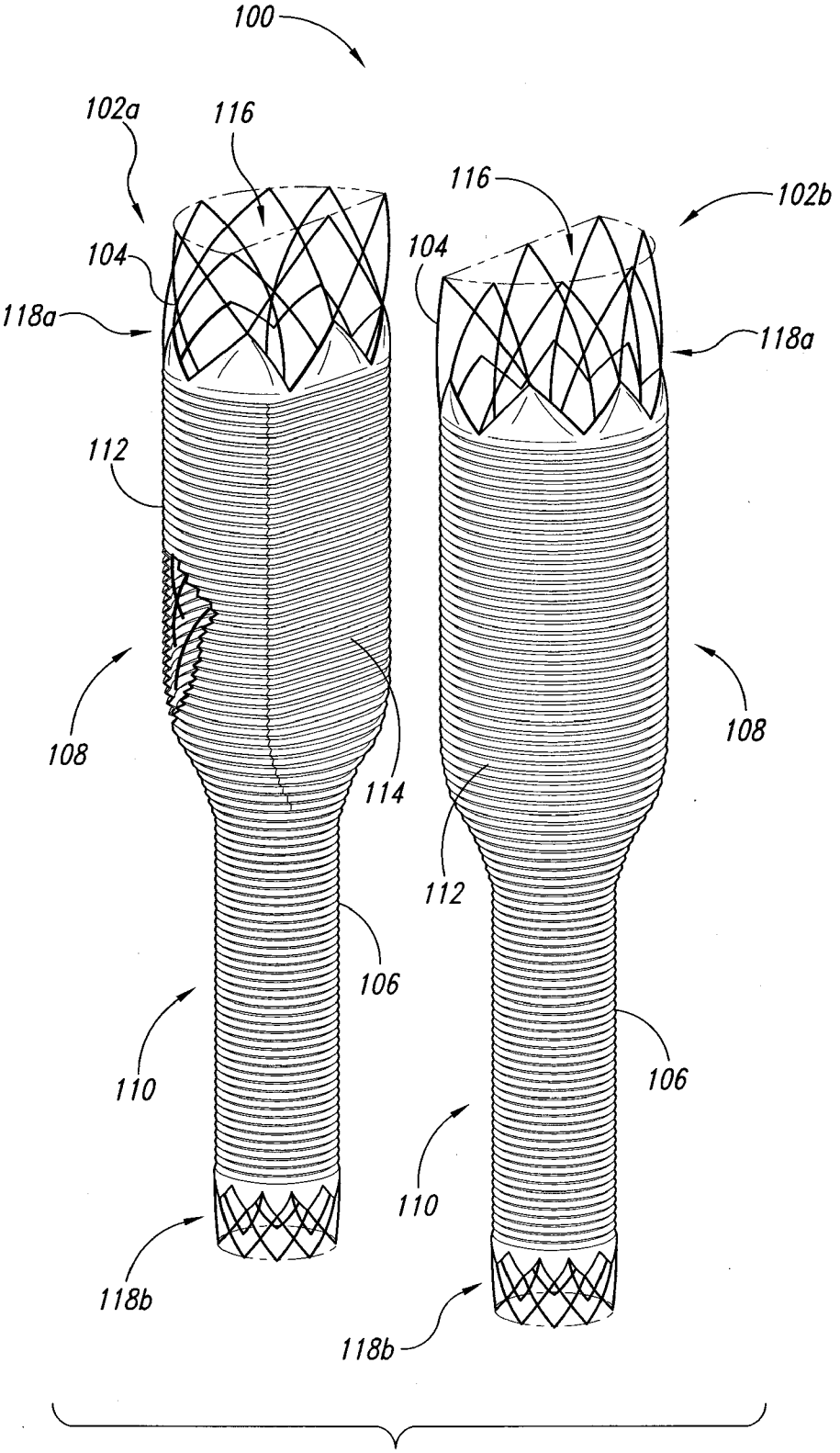

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Fig. 1A

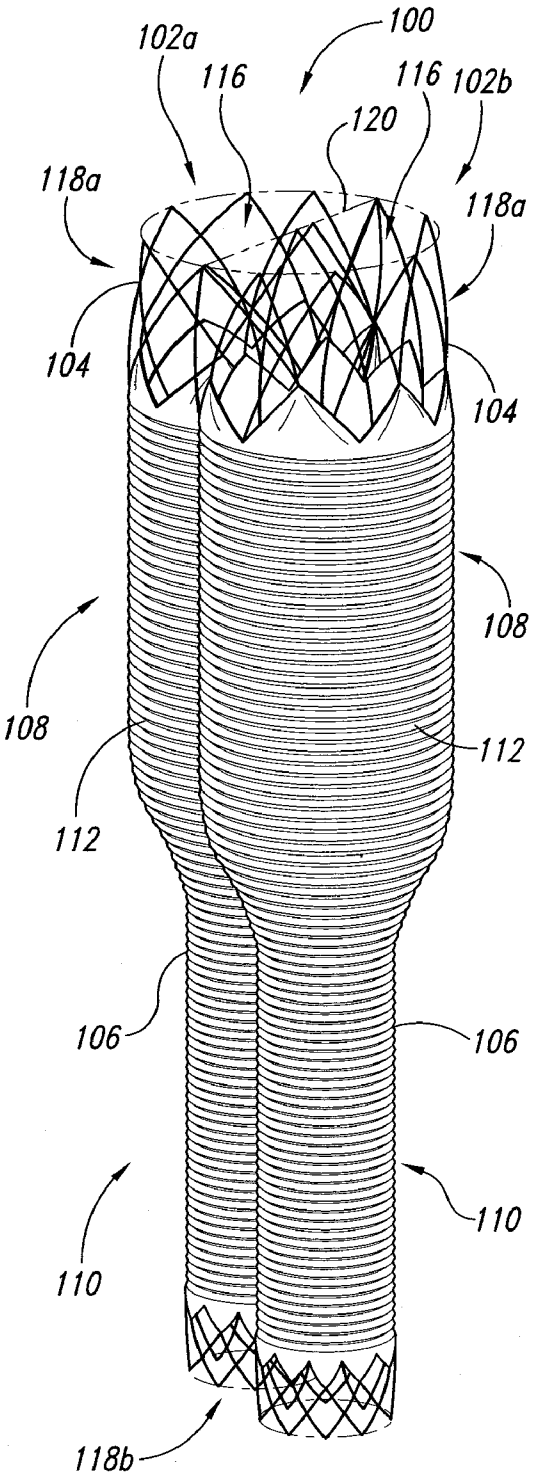
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Fig. 1B

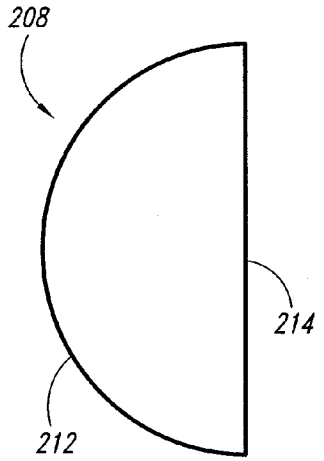


Fig. 2A

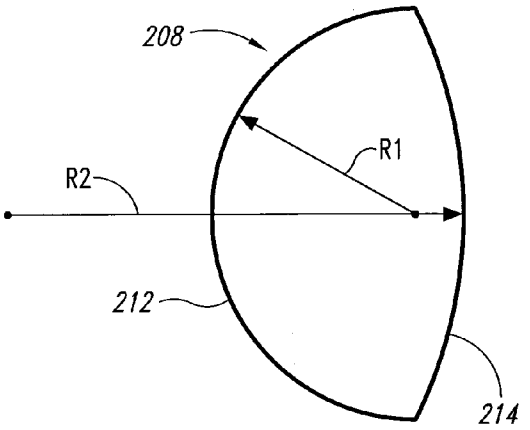


Fig. 2B

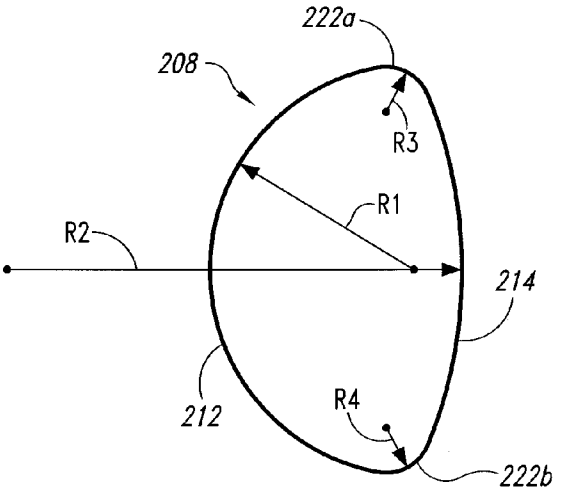
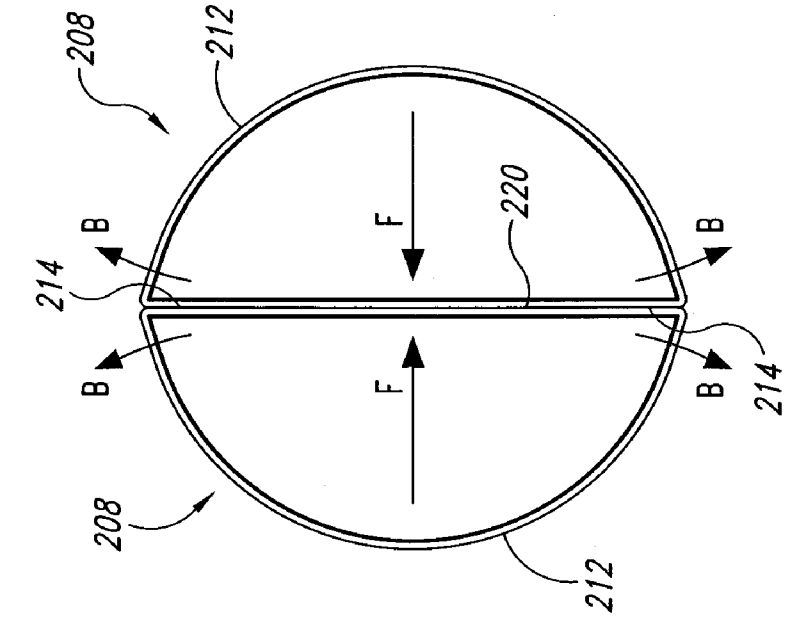


Fig. 2C



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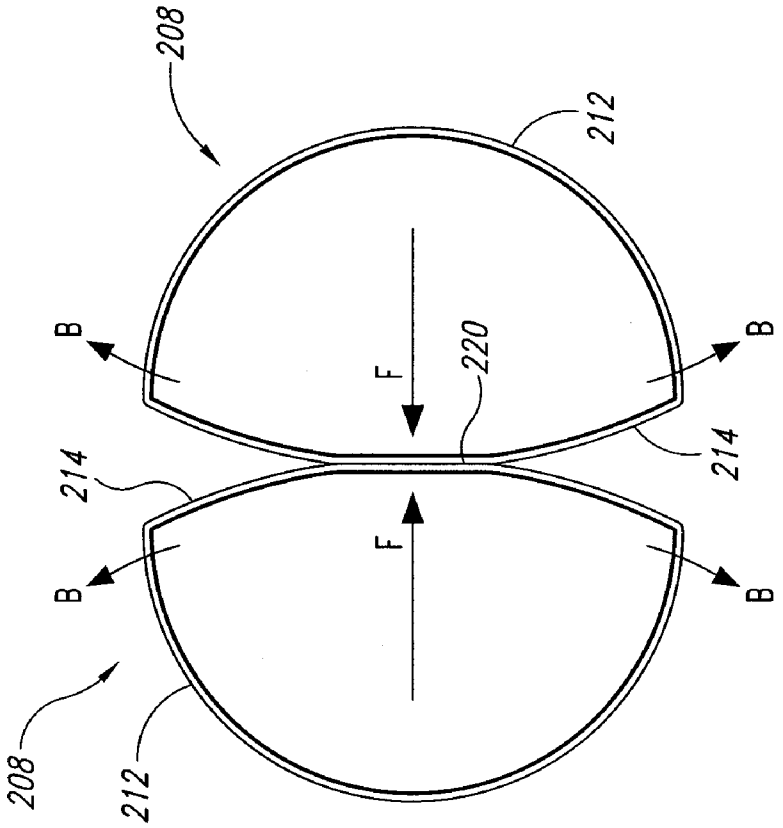

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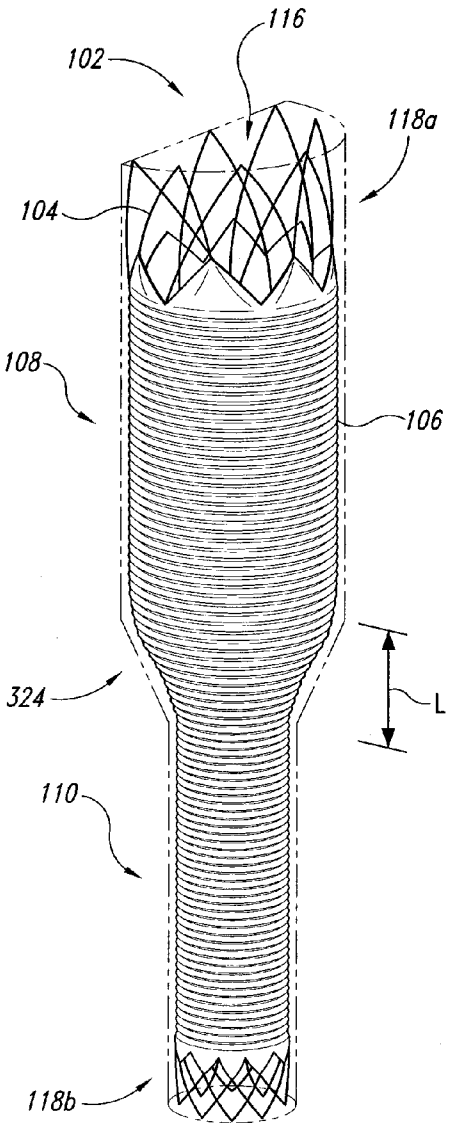
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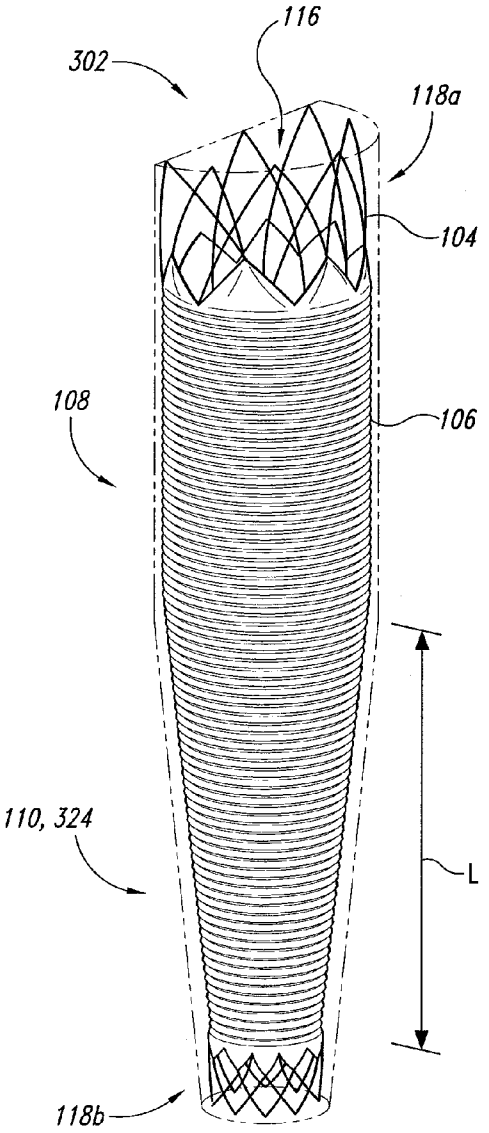
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Fig. 3B

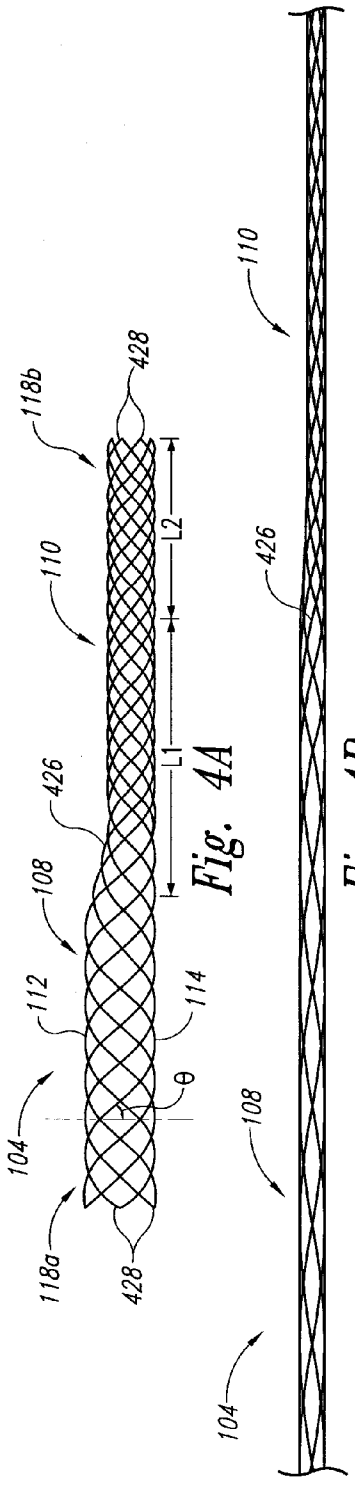
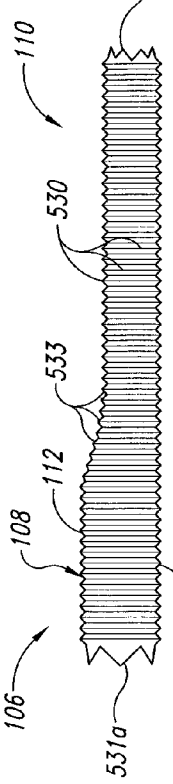


Fig. 4A

Fig. 4B



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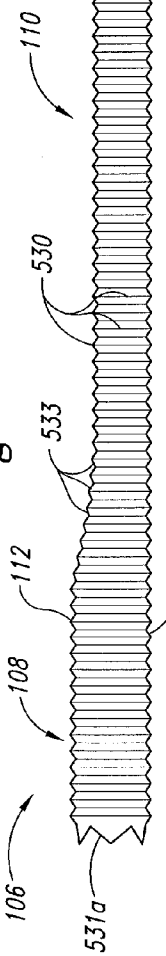

Fig. 5B

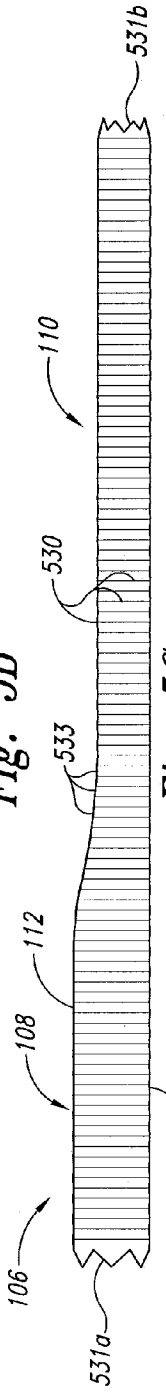

Fig. 5C

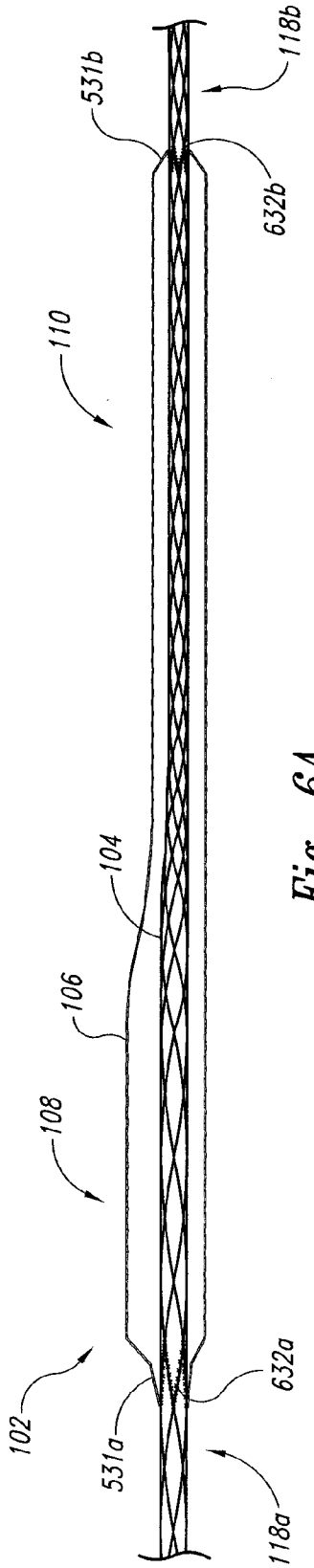
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Fig. 6A

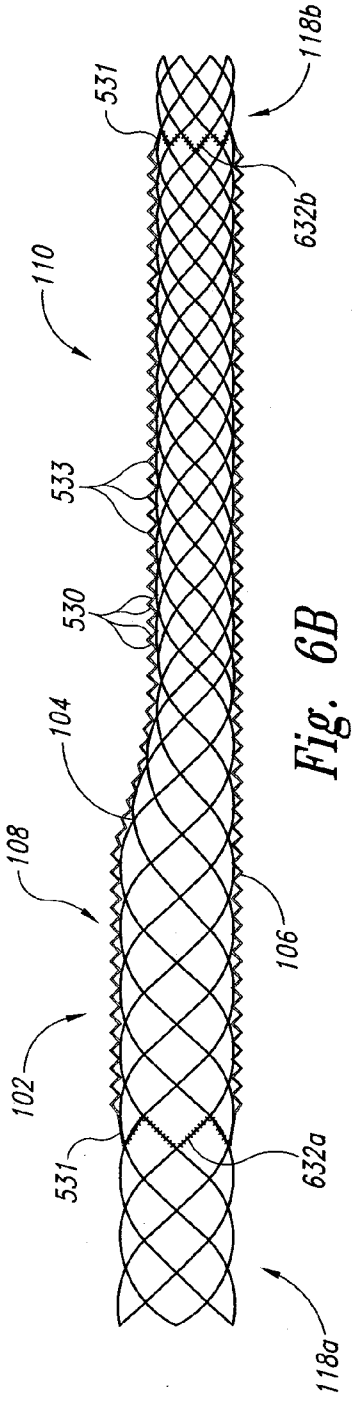
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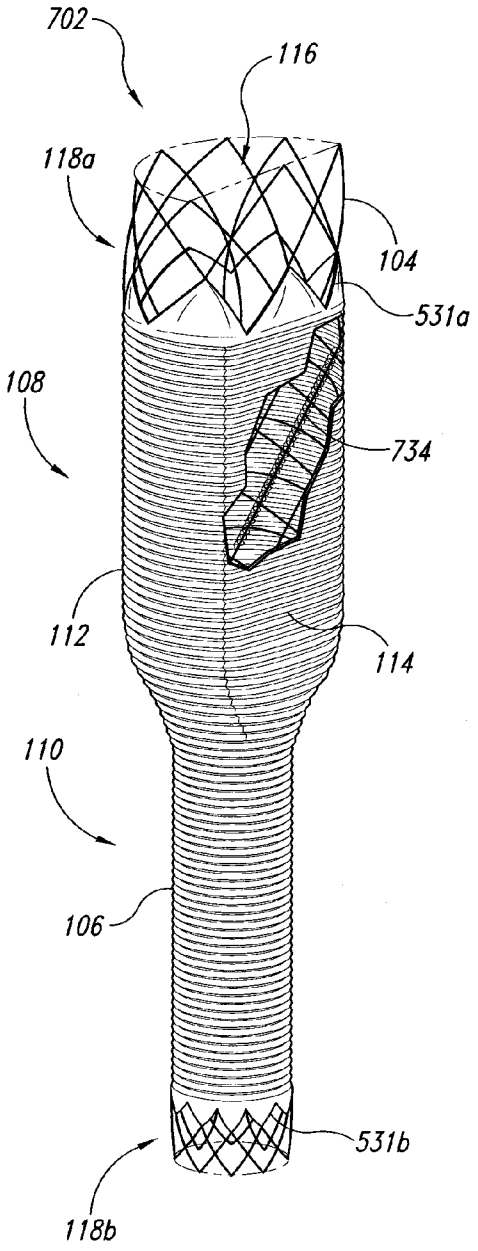
Fig. 6B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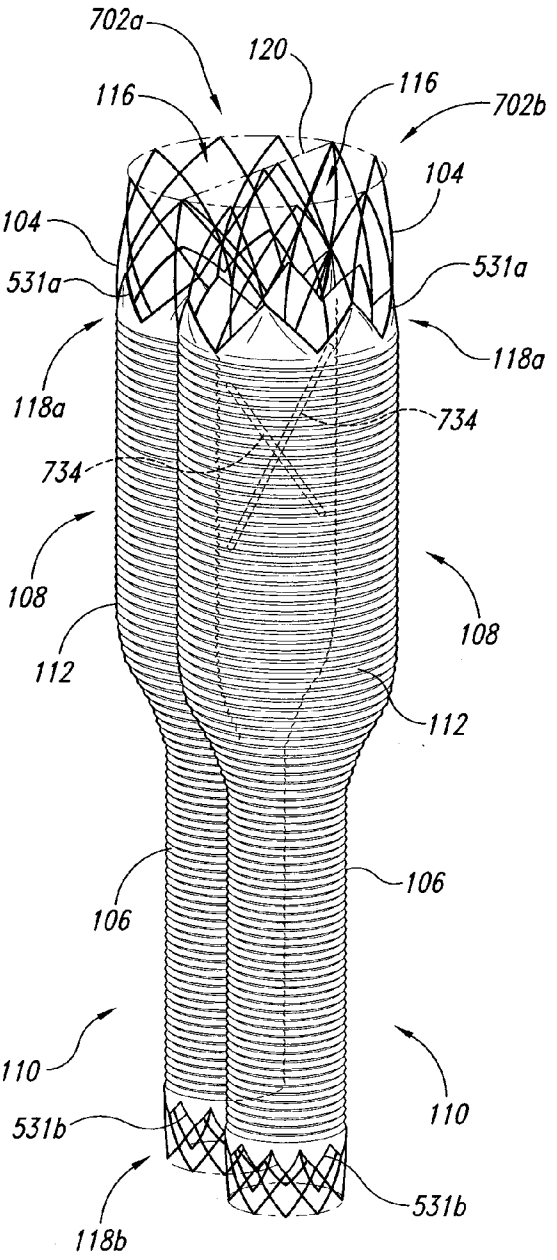
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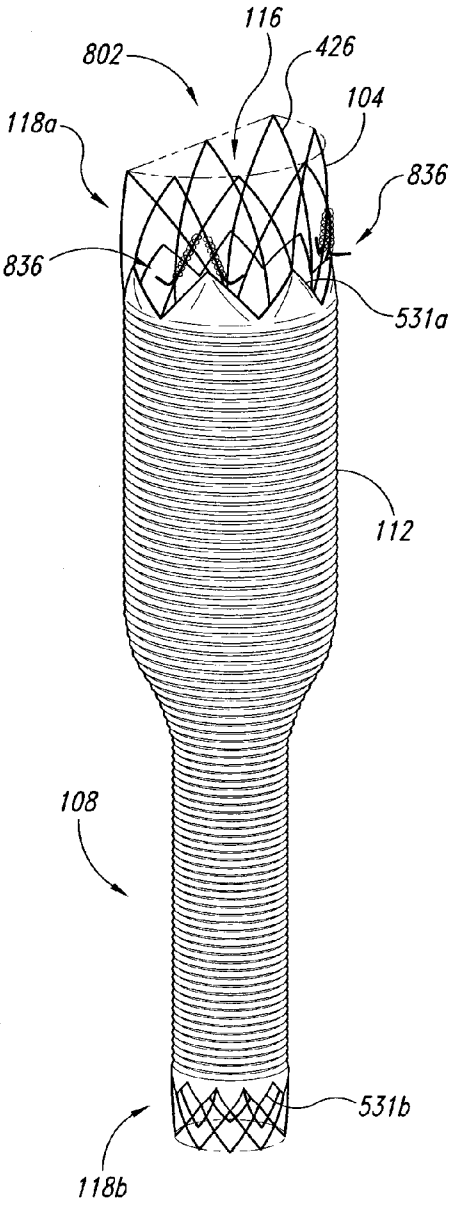
Fig. 7B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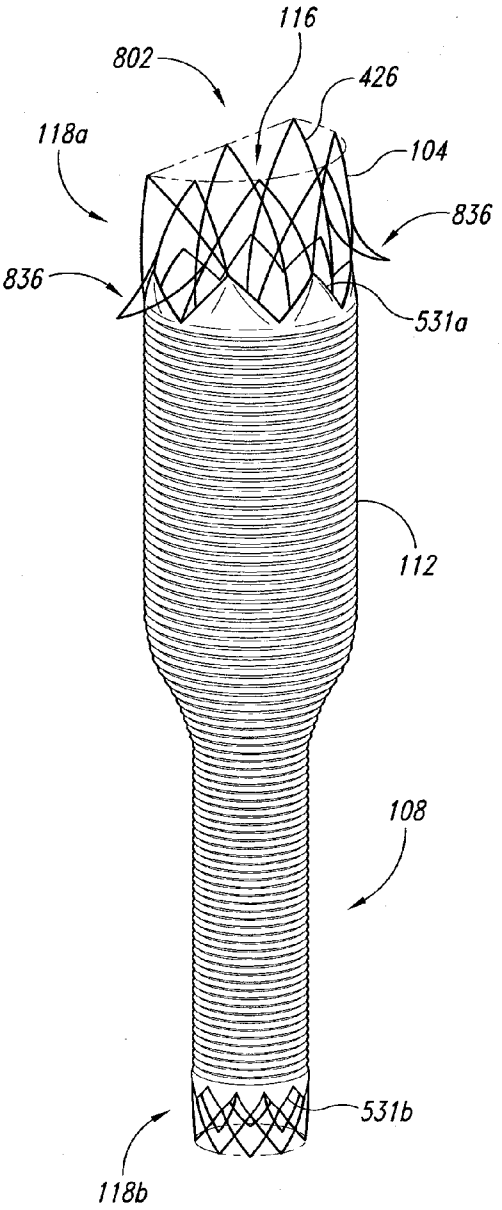
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Fig. 8B

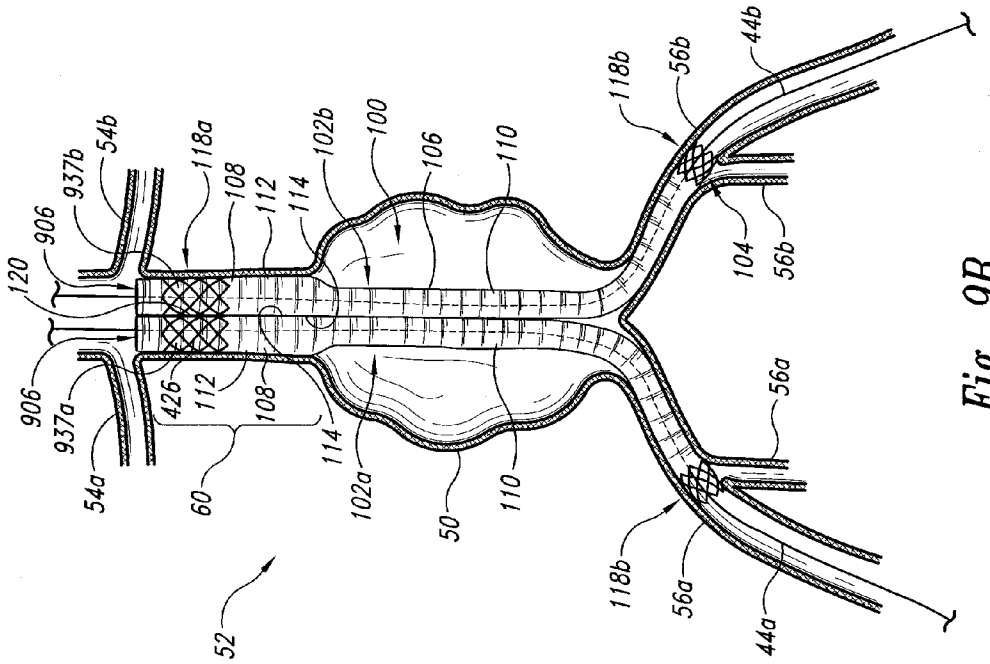
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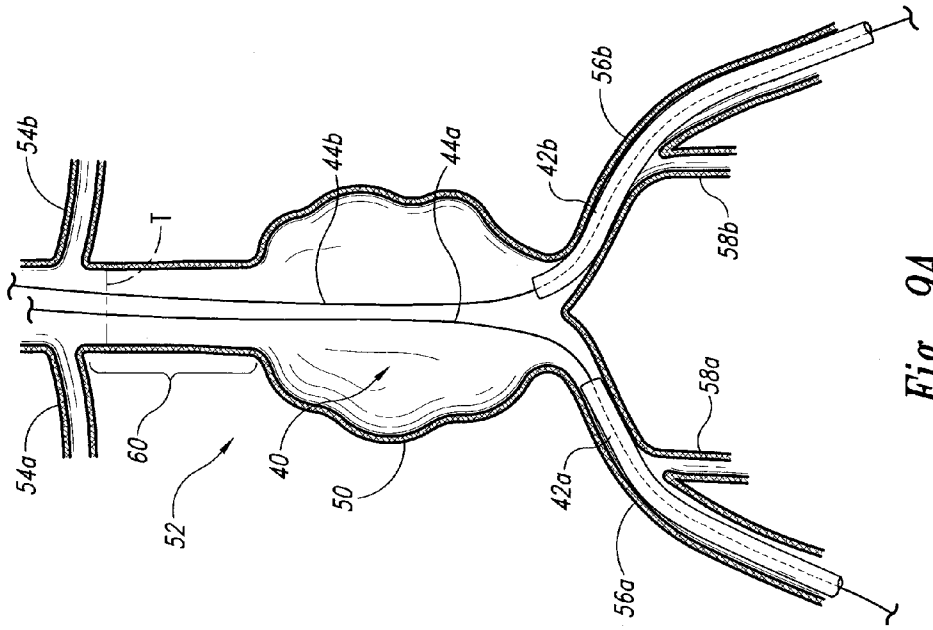
Fig. 9B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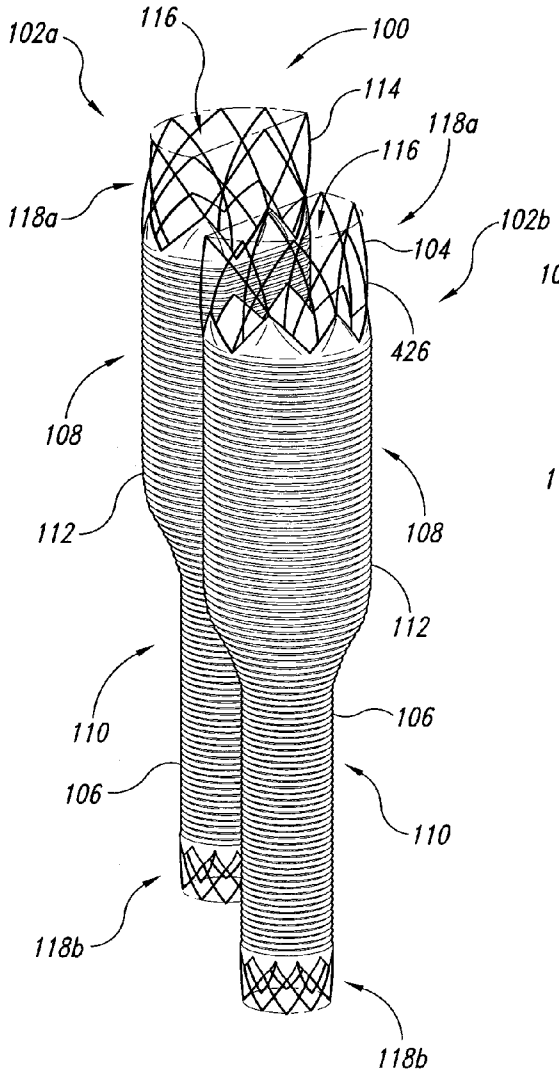

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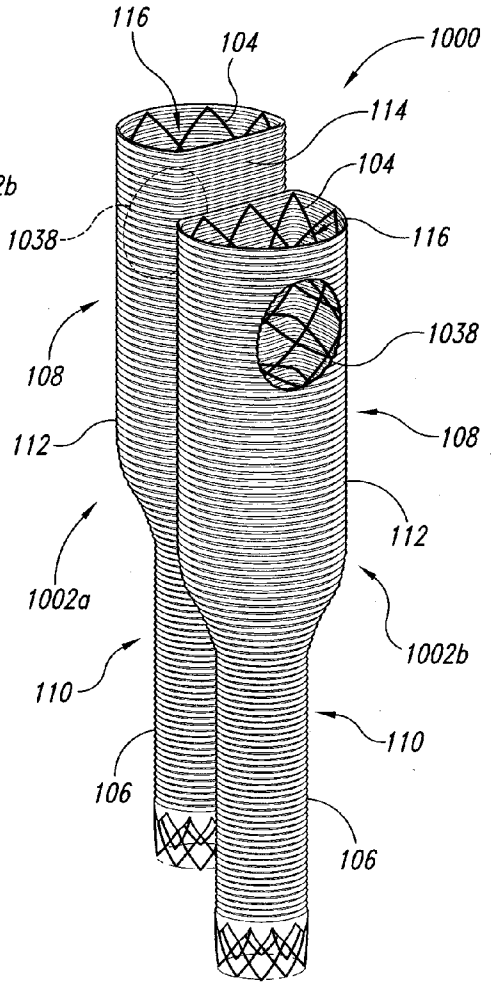


Fig. 10B

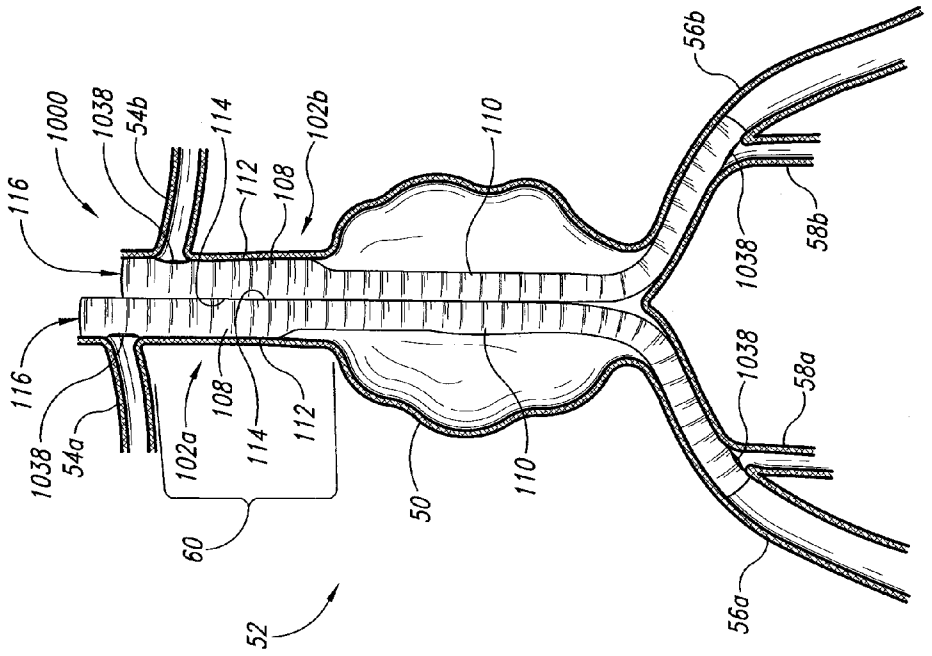


Fig. 11B

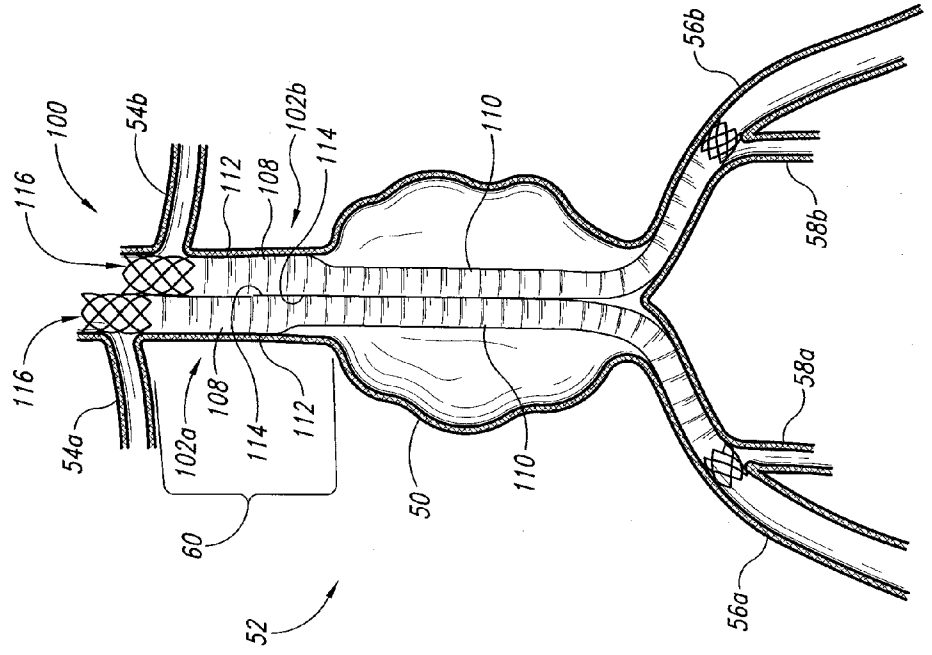
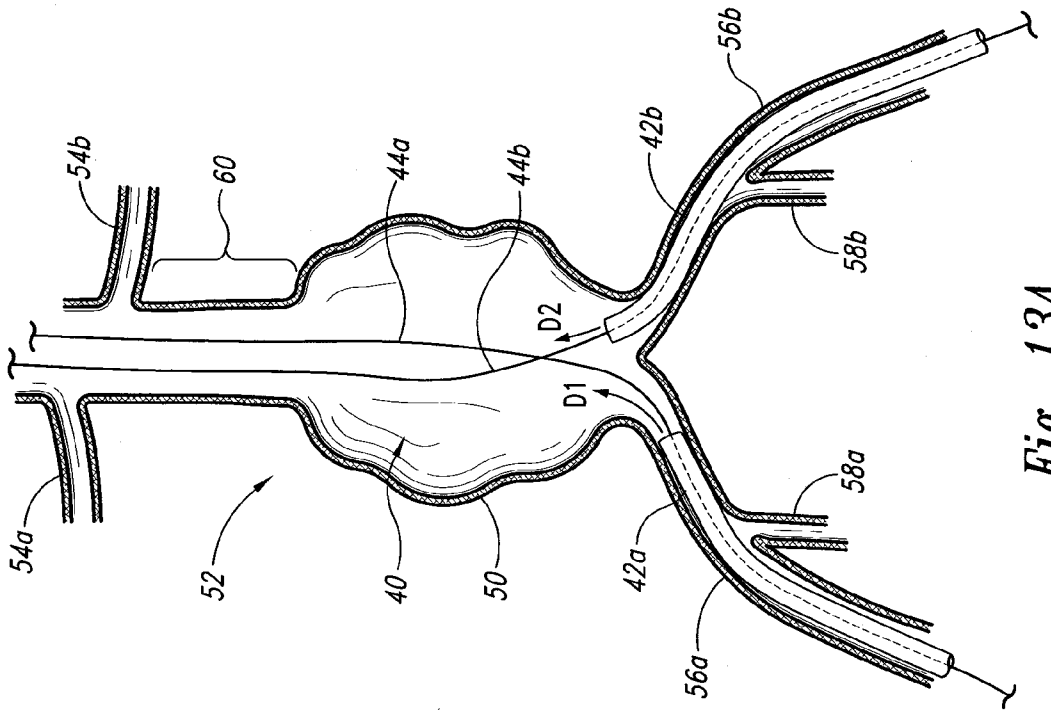


Fig. 11A



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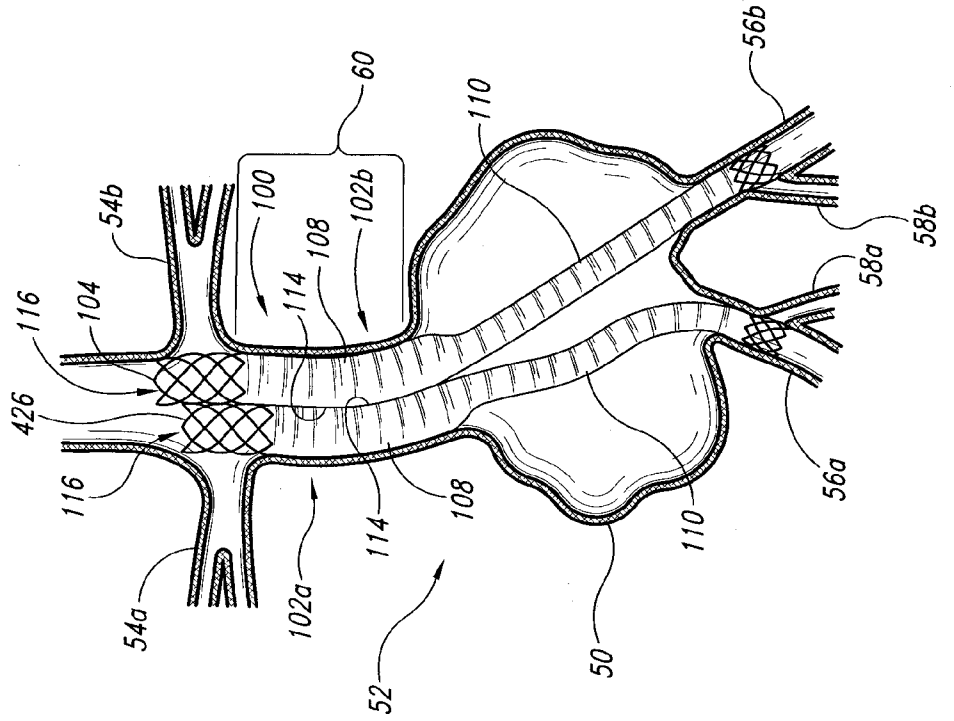


Fig. 12

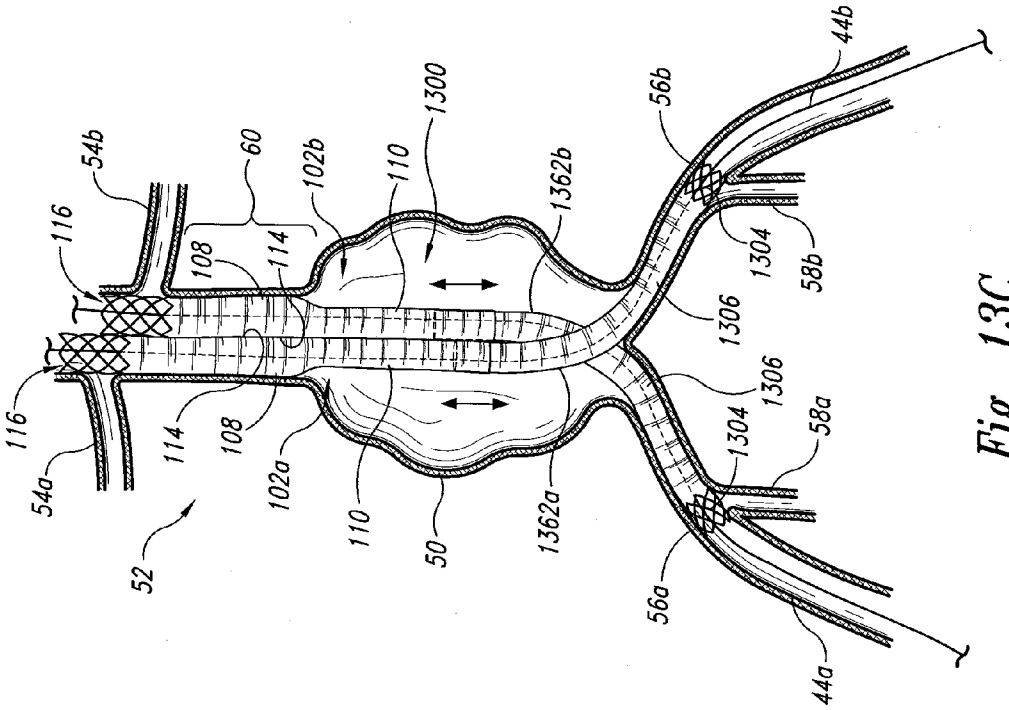


Fig. 13C

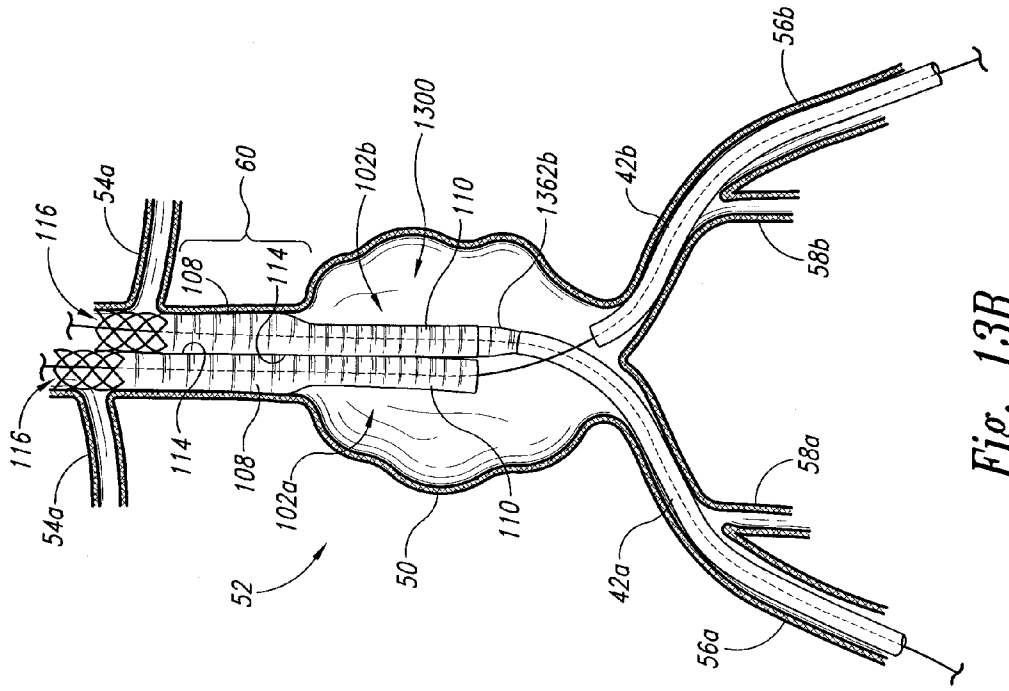


Fig. 13B

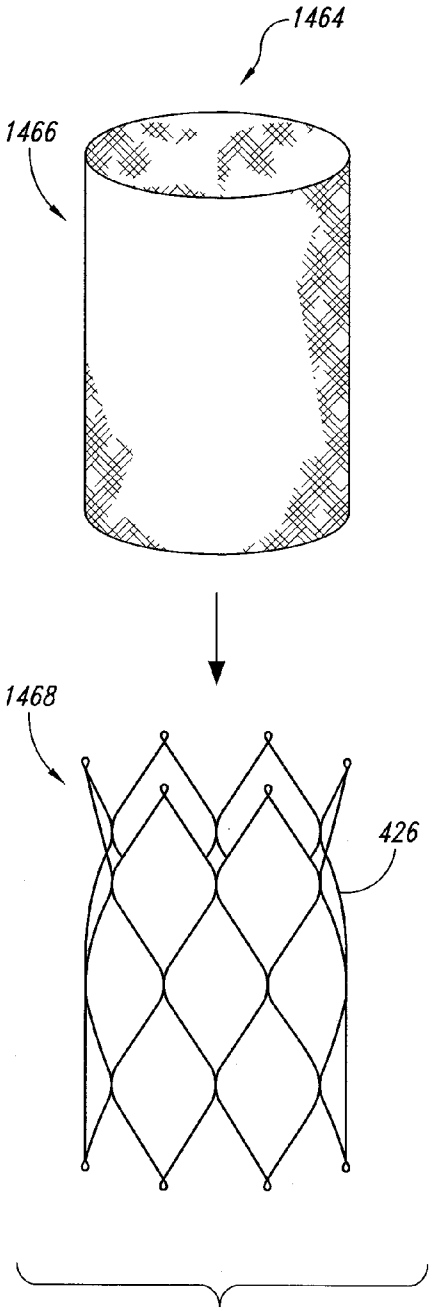
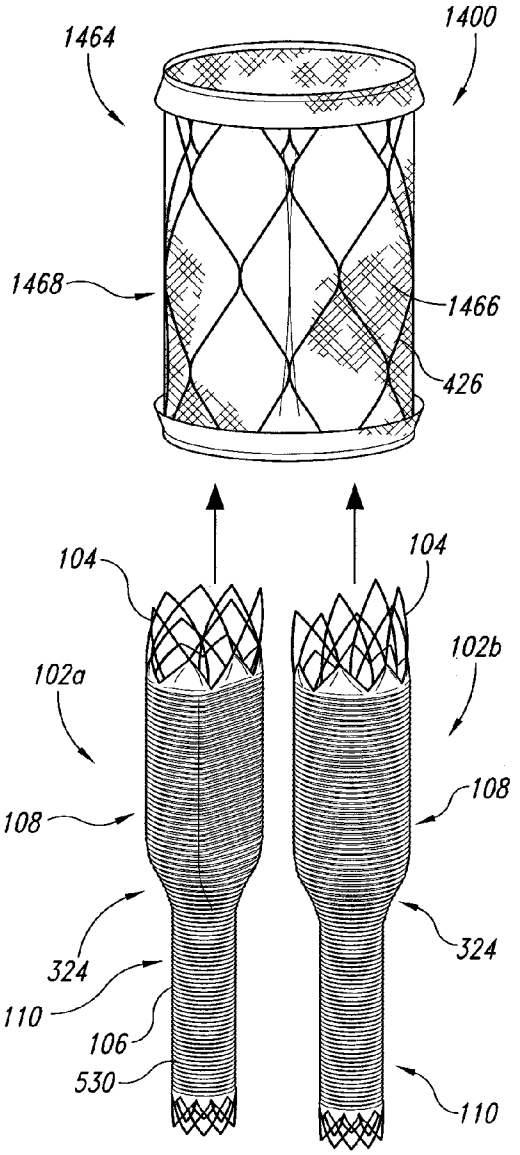


Fig. 14A



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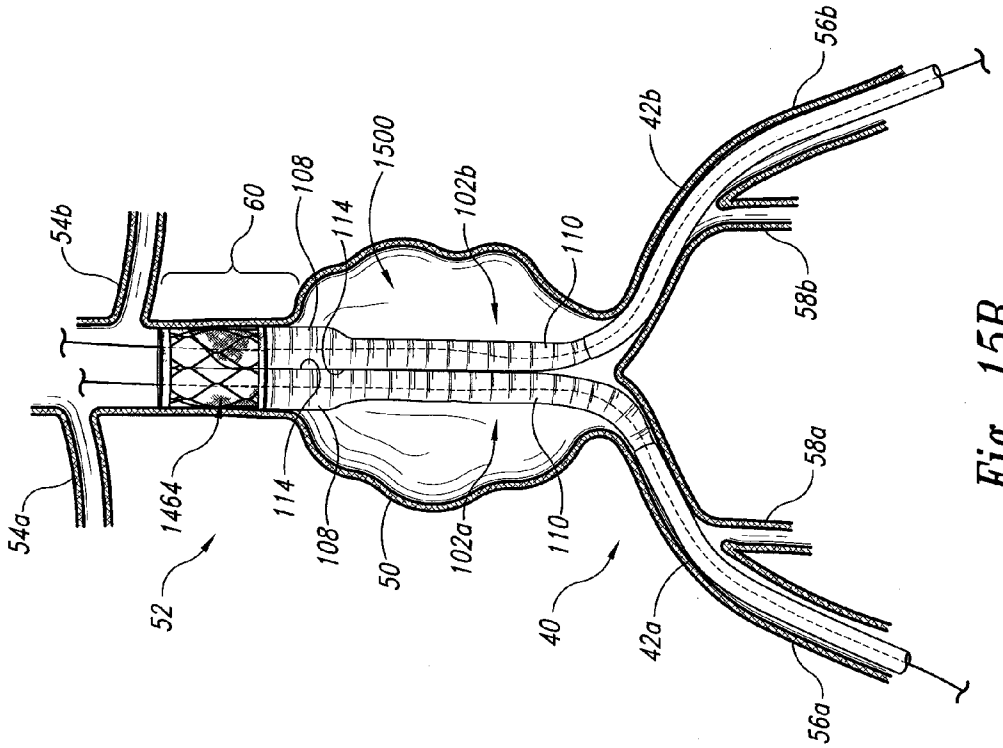

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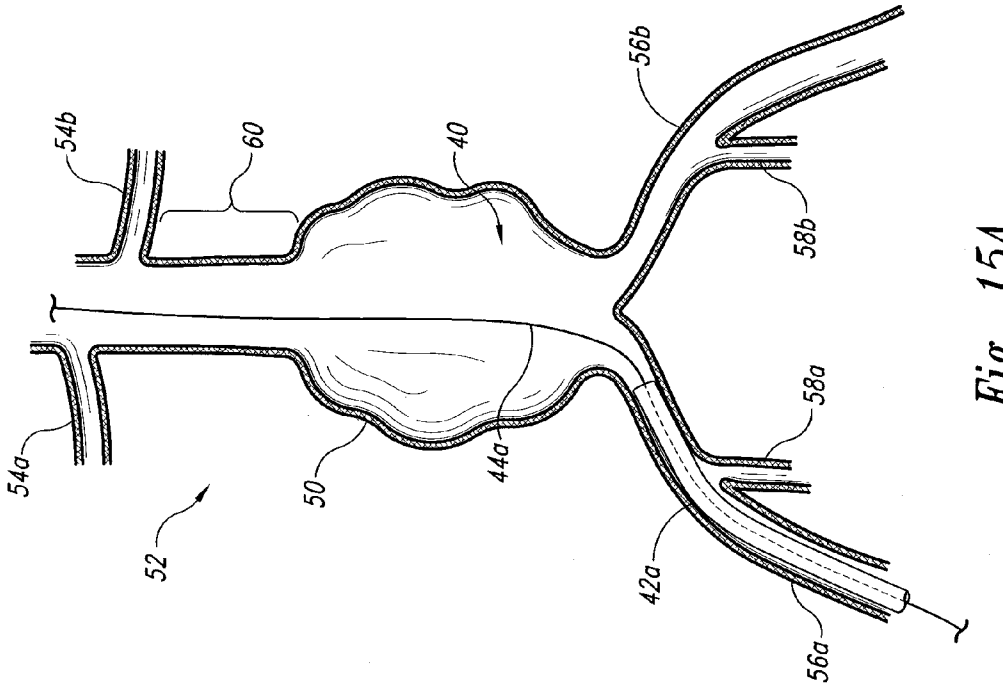

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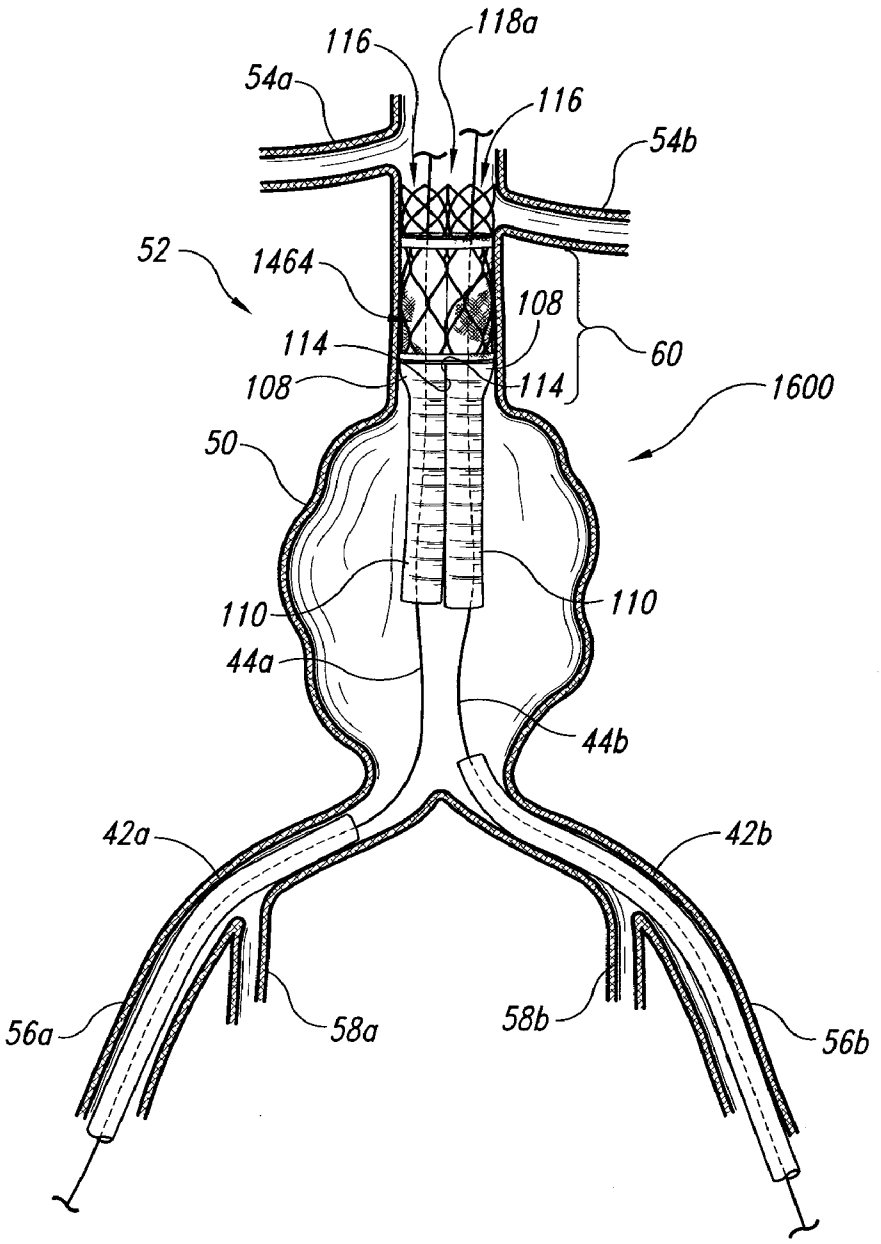
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Fig. 16

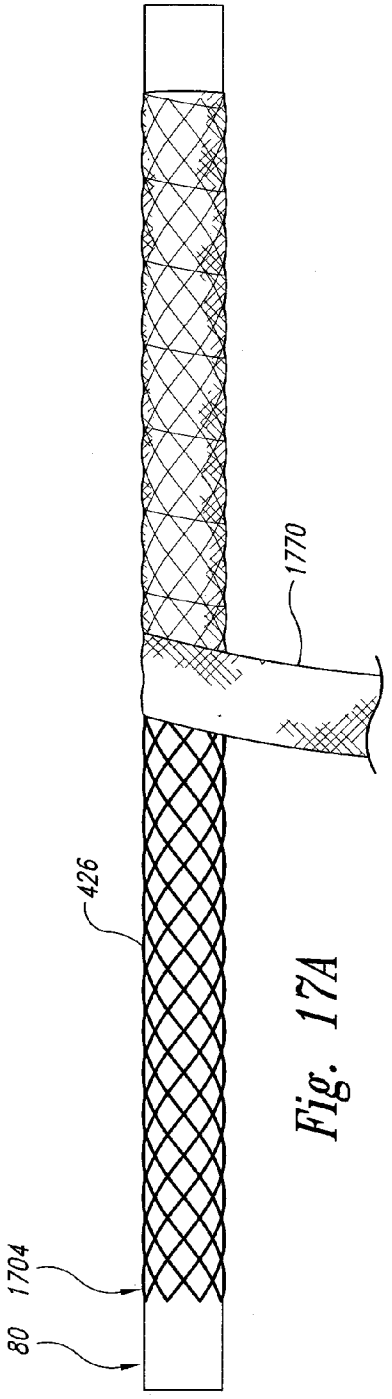

Fig. 17A

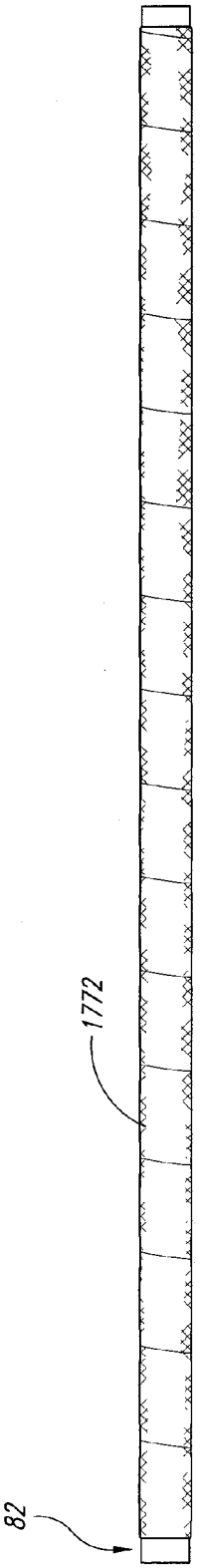

Fig. 17B

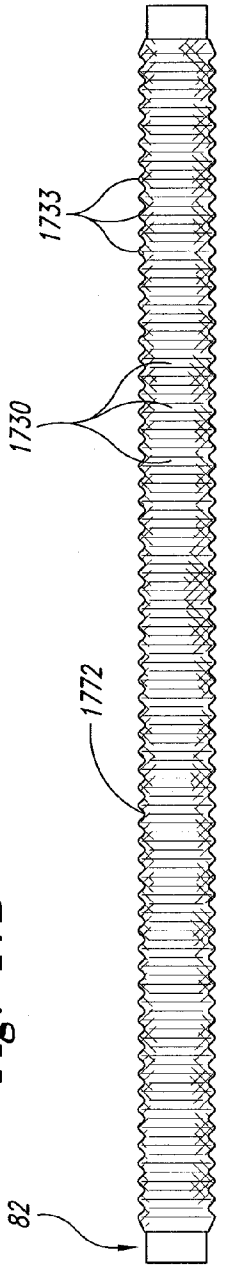

Fig. 17C

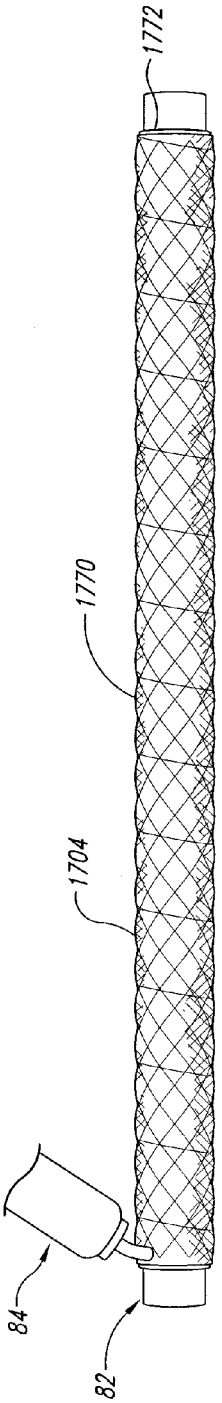

Fig. 17D

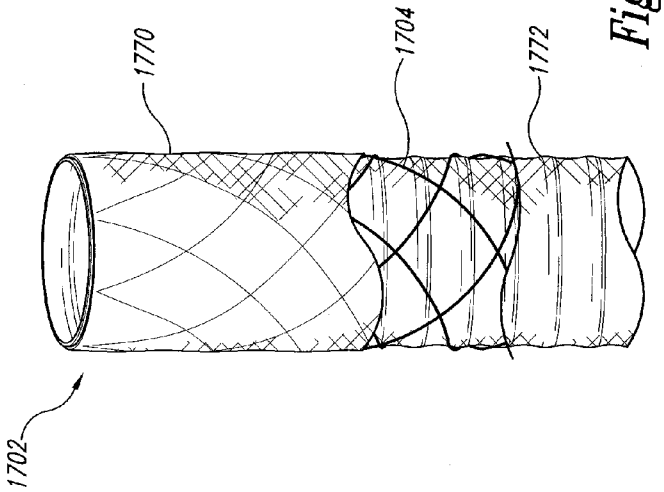

Fig. 17E

MODULAR ENDOGRAFT DEVICES AND ASSOCIATED SYSTEMS AND METHODS

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] The present application claims priority to each of the following U.S. Provisional Applications:

[0002] (A) U.S. Provisional Application No. 61/265,713, filed on Dec. 1, 2009, entitled "IMPROVED SYSTEMS AND METHODS FOR MODULAR ABDOMINAL AORTIC ANEURYSM GRAFT;" and

[0003] (B) U.S. Provisional Application No. 61/293,581, filed Jan. 11, 2010, entitled "IMPROVED SYSTEMS AND METHODS FOR MODULAR ABDOMINAL AORTIC ANEURYSM GRAFT."

[0004] All of the foregoing applications are incorporated herein by reference in their entireties.

TECHNICAL FIELD

[0005] The present technology generally relates to endograft devices and methods for percutaneous endovascular delivery of the endograft devices across aneurysms. In particular, several embodiments are directed toward a modular bi-luminal endograft device with independently positioned components for endovascular aneurysm repair.

BACKGROUND

[0006] An aneurysm is a dilation of a blood vessel at least 1.5 times above its normal diameter. The dilated vessel can form a bulge known as an aneurysmal sac that can weaken vessel walls and eventually rupture. Aneurysms are most common in the arteries at the base of the brain (i.e., the Circle of Willis) and in the largest artery in the human body, the aorta. The abdominal aorta, spanning from the diaphragm to the aortoiliac bifurcation, is the most common site for aortic aneurysms. The frequency of abdominal aortic aneurysms ("AAAs") results at least in part from decreased levels of elastins in the arterial walls of the abdominal aorta and increased pressure due to limited transverse blood flow.

[0007] Aneurysms are often repaired using open surgical procedures. Surgical methods for repairing AAAs, for example, require opening the abdominal region from the breast bone to the pelvic bone, clamping the aorta to control bleeding, dissecting the aorta to remove the aneurysmal section, and attaching a prosthetic graft to replace the diseased artery. The risks related to general anesthesia, bleeding, and infection in these types of open surgical repairs result in a high possibility of operative mortality. Thus, surgical repair is not a viable option for many patients. Moreover, the recovery process is extensive for the patients fit for surgical repair. An open surgical repair of an AAA generally requires seven days of post-operational hospitalization and, for uncomplicated operations, at least six to eight weeks of recovery time. Thus, it is a highly invasive and expensive procedure.

[0008] Minimally invasive surgical techniques that implant prosthetic grafts across aneurysmal regions of the aorta have been developed as an alternative or improvement to open surgery. Endovascular aortic repairs ("EVAR"), for example, generally require accessing an artery (e.g., the femoral artery) percutaneously or through surgical cut down, introducing guidewires into the artery, loading an endograft device into a catheter, and inserting the loaded catheter in the artery. With the aid of imaging systems (e.g., X-rays), the endograft device can be guided through the arteries and deployed from a distal opening of the catheter at a position superior to the aneurysm. From there, the endograft device can be deployed across the aneurysm such that blood flows through the endograft device and bypasses the aneurysm.

[0009] EVAR devices should be implanted at a precise location across the aneurysmal region and securely fixed to the vessel wall because improper placement, migration, and/or projection of the endograft device into branching vessels may interfere with the blood flow to nearby physiological structures. For example, to avoid impairing renal functions, the endograft device should not inhibit blood flow to the renal arteries. In addition to the variations in the vasculature between patients, the characteristics of the aneurysms themselves can also pose challenges because of the anatomical variations and the different structural features of individual aneurysms. For example, the vascular bifurcation at the iliac arteries and the angulation of aneurysmal sacs are both known to pose challenges to methods and devices for treating AAAs. Conventional systems address these challenges by having many different EVAR devices with different sizes and shapes.

BRIEF DESCRIPTION OF THE DRAWINGS

[0010] FIG. **1**A is a partial cut-away, isometric view of a modular endograft system configured in accordance with an embodiment of the technology.

[0011] FIG. **1**B is an isometric view of the modular endograft system of FIG. **1**A configured in accordance with an embodiment of the technology.

[0012] FIGS. **2**A-C are cross-sectional top views of superior portions for endograft devices shaped in accordance with embodiments of the technology.

[0013] FIGS. **2**D and **2**E are cross-sectional top views of the superior portion of FIG. **2**B being mated with a complementary superior portion in accordance with an embodiment of the technology.

[0014] FIGS. **3**A and **3**B are isometric views of endograft devices configured in accordance with embodiments of the technology.

[0015] FIGS. **4**A and **4**B are side views of an integrated frame in an expanded configuration and in a low-profile configuration, respectively, in accordance with an embodiment of the technology.

[0016] FIGS. **5**A-C are side views of a cover being extended from an expanded configuration to a low-profile configuration in accordance with an embodiment of the technology.

[0017] FIGS. **6**A and **6**B are cross-sectional views of an endograft device in a low-profile configuration and in an expanded configuration, respectively, in accordance with embodiments of the technology.

[0018] FIGS. **7**A and **7**B are isometric views of endograft devices configured in accordance with other embodiments of the technology.

[0019] FIGS. **8**A and **8**B are isometric views of endograft devices configured in accordance with further embodiments of the technology.

[0020] FIGS. **9**A and **9**B are schematic views of a two-part modular endograft system being deployed across an aneurysm in accordance with an embodiment of the technology.

[0021] FIGS. **10**A and **10**B are isometric views of modular endograft systems configured in accordance with additional embodiments of the technology.

[0022] FIGS. **11**A and **11**B are schematic views of the modular endograft system of FIG. **10**A and the modular endograft system of FIG. **10**B, respectively, deployed across aneurysms in accordance with other embodiments of the technology.

[0023] FIG. **12** is a schematic view of the modular endograft system of FIG. **9**B deployed across an aneurysm in accordance with a further embodiment of the technology.

[0024] FIGS. **13**A-C are schematic views of a four-part modular endograft system being deployed across an aneurysm in accordance with an embodiment of the technology.

[0025] FIGS. **14**A and **14**B are isometric views of a modular endograft system configured in accordance with an additional embodiment of the technology.

[0026] FIGS. **15**A and **15**B are schematic views of a three-part modular endograft system being deployed across an aneurysm in accordance with an embodiment of the technology.

[0027] FIG. **16** is a schematic view of a five-part modular endograft system being deployed across an aneurysm in accordance with an embodiment of the technology.

[0028] FIGS. **17**A-E are views of coating layers being applied to an integrated frame in accordance with an embodiment of the technology.

DETAILED DESCRIPTION

[0029] Specific details of several embodiments of the technology are described below with reference to FIGS. **1**A-**17**E. Although many of the embodiments are described below with respect to devices that at least partially repair abdominal aortic aneurysms ("AAAs"), other applications and other embodiments are within the scope of the technology. For example, the technology can be used to repair aneurysms in other portions of the vasculature. Additionally, several other embodiments of the technology can have different configurations, components, or procedures than those described in this section. A person of ordinary skill in the art, therefore, will accordingly understand that the technology may have other embodiments with additional elements, or the technology may have other embodiments without several of the features shown and described below with reference to FIGS. **1**A-**17**E.

[0030] With regard the use of "superior" and "inferior" within this application, inferior generally refers being situated below or directed downward, and superior generally refers to being situated above or directed upward.

[0031] With regard to the use of "expansion" and "constriction" within this application, expansion refers to a radial increase in a cross-sectional dimension of a device or component, and constriction refers to a radial decrease in the cross-sectional dimension of the device or component. For example, FIG. **4**A shows an integrated frame **104** in an expanded configuration, and FIG. **4**B shows the integrated frame **104** in a constricted configuration.

[0032] With regard to the use of "contraction" and "extension" within this application, contraction refers to a longitudinal decrease in the length of a device or component, and extension refers to a longitudinal increase in the length of the device or component. For example, FIG. **5**A shows a cover **106** in a contracted configuration, and FIG. **5**C shows the cover **106** in an extended configuration.

[0033] With regard to the terms "distal" and "proximal" within this application, the terms can reference a relative position of the portions of an implantable device and/or a delivery device with reference to an operator. Proximal refers to a position closer to the operator of the device, and distal refers to a position that is more distant from the operator of the device.

[0034] 1. Endograft System Structures

[0035] 1.1 Selected Endograft Devices

[0036] FIGS. **1**A and **1**B are isometric views of a modular endograft system **100** ("system **100**") in accordance with an embodiment of the technology. The system **100** can include separate endograft devices **102** (identified individually as a first endograft device **102***a* and a second endograft device **102***b*) that can be coupled, mated, or otherwise substantially sealed together in situ. Each endograft device **102**, for example, can include an integrated frame **104** ("frame **104**") and a substantially impermeable cover **106** ("cover **106**") extending over at least a portion of the frame **104**. The frame **104** and the cover **106** of an individual endograft device **102** can form a discrete lumen **116** through which blood can flow to bypass an aneurysm. In operation, the endograft devices **102** are generally delivered separately and positioned independently across the aneurysm.

[0037] As shown in FIGS. **1**A and **1**B, each endograft device **102** includes a superior portion **108** and an inferior portion **110**. The superior portion **108** can include a convexly curved outer wall **112** and a septal wall **114**. As shown in FIG. **1**A, the septal wall **114** can be substantially flat such that the superior portion **108** forms a "D" shape at a superior portion of the lumen **116**. In other embodiments, the septal wall **114** can be convexly curved with a larger radius of curvature than the outer wall **112** such that the superior portion **108** forms a complex ellipsoid having another D-shaped cross-section at the superior portion of the lumen **116**. In further embodiments, the superior portion **108** can have asymmetrical shapes or other suitable cross-sectional configurations that can mate with each other in the septal region and mate with an arterial wall around the periphery of the outer wall **112**. The inferior portion **110** can have a circular cross-sectional shape as illustrated in FIG. **1**A, or the inferior portion **110** can have an elliptical shape, a rectangular shape, an asymmetrical shape, and/or another suitable cross-sectional shape for an inferior portion of the lumen **116**.

[0038] The superior portions **108** of the endograft devices **102** are mated together and at least substantially sealed along the septal walls **114** within the aorta above the aneurysm. In some embodiments, the superior portion **108** can be approximately 2-4 cm in length to adequately fix the outer walls **112** to the arterial walls such that they are at least substantially sealed together. In other embodiments, the superior portion **108** can be longer or shorter. In one embodiment in accordance with the technology, the inferior portions **110** can extend through an inferior portion of the aneurysm and into corresponding iliac arteries to bypass the aneurysm. In another embodiment, one or both inferior portions **110** can terminate within the aneurysm to form what is known to those skilled in the art as a "gate." As described in further detail below, limbs (not shown) can be attached to the proximal ends of the inferior portions **110** and extended into the iliac arteries to bypass the aneurysm.

[0039] In the embodiment shown in FIGS. **1**A and **1**B, the frames **104** have bare end portions **118** (identified individually as first end portions **118***a* and second end portions **118***b*) that extend beyond the covers **106**. As shown in FIGS. **1**A and **1**B, the first end portion **118***a* can extend distally from the superior terminus of the cover **106**, and the second end por-

tion **118***b* can extend proximally from the inferior terminus of the cover **106**. In some embodiments, the end portions **118** can be trumpeted or flared to interface with the arterial walls of the aorta and/or the iliac arteries. This can promote cell ingrowth that strengthens the seal between the endograft devices **102** and the adjacent arteries.

[0040] The end portions **118** can also increase the available structure for securing the endograft device **102** to the artery and increase the surface area of the covers **106** for sealably fixing the endograft devices **102** to arterial walls. This decreases the precision necessary to position the endograft devices **102** and increases the reliability of the implanted system **100**. For example, a short infrarenal aortic neck (e.g., less than 2 cm) generally requires precise placement of the endograft devices **102** to preserve blood flow to the renal arteries while still providing enough surface area for the endograft devices **102** to be properly affixed with the aorta. In the embodiment shown in FIGS. **1**A and **1**B, however, the first end portions **118***a* can be placed at the entrance of the renal arteries to allow lateral blood flow into the renal arteries and provide a larger structure for fixing the endograft devices **102** to the arterial wall and a larger sealing area with the arterial wall. The end portions **118** can also provide accessible sites for recapture (e.g., by guidewires, bead and collet, etc.) that enhance the accuracy of positioning the endograft devices **102** across the aneurysm.

[0041] During deployment of the system **100**, each endograft device **102** can be delivered independently to an aneurysmal region in a low-profile configuration. The low-profile configuration has a first cross-sectional dimension and a first length that can facilitate percutaneous endovascular delivery of the system **100**. Because each device **102** extends around only a portion of the vessel periphery, the individual endograft devices **102** can be constricted (i.e., radially collapsed) to a smaller diameter than conventional AAA devices with a single superior portion that extends around the complete periphery of the vessel wall. In some embodiments, for example, each of the endograft devices **102** can have a diameter of 25 mm in the expanded configuration, and can be constricted to a diameter of 4 mm in the low-profile configuration to be percutaneously deployed across the aneurysm through a 12 F catheter. Additionally, as described in more detail below, because each endograft device **102** is delivered independently, the end portions **118** and fenestrations can facilitate staggering the endograft devices **102** to accommodate asymmetrical anatomies.

[0042] At a target site in the aneurysmal region, the endograft devices **102** can self-expand to an expanded configuration (e.g., shown in FIGS. **1**A and **1**B). The expanded configuration can have a second cross-sectional dimension greater than the first cross-sectional dimension and a second length less than the first length. In the expanded configuration shown in FIG. **1**B, the septal wall **114** (FIG. **1**A) of the first endograft device **102***a* can be forced against the opposing septal wall **114** of the second endograft device **102***b*. When in situ within the aorta, the forces between the opposing septal walls **114** form a septum **120** in which the first and second septal walls **114** are at least substantially sealed together to prevent blood from flowing between the endograft devices **102** and into the aneurysm. Additionally, as shown in FIG. **1**B, the texture (e.g., ribbing) on the covers **106** can mate at the septum **120** to further strengthen the seal between the septal walls **114**. Similarly, the texture of the cover **106** on the outer walls **112** can interface with the adjacent vessel walls to strengthen the seal around the periphery of the endograft devices **102**.

[0043] In operation, the system **100** can prevent blood from collecting in a diseased aneurysmal portion of a blood vessel (e.g., the aorta, the iliac arteries, etc.). Rather, the system **100** can direct blood into the lumens **116**, funnel the blood through the superior and inferior portions **108** and **110**, and discharge the blood into healthy portions of the iliac arteries, thereby at least substantially bypassing the aneurysm. The bifurcated system **100** facilitates independent positioning of the first and second endograft devices **102** to accommodate disparate structures and morphologies of the abdominal aorta and/or iliac arteries. For example, the first endograft device **102***a* can be positioned independently in a desired location without being constrained by a desired placement of the second endograft device **102***b*. Accordingly, the system **100** can easily adapt to a variety of different anatomies and thereby provide a modular alternative to customized endograft systems.

[0044] 1.2 Select Embodiments of Superior Portions

[0045] FIGS. **2**A-C are cross-sectional top views of superior portions **208** of endograft devices (e.g., endograft devices **102** shown in FIGS. **1**A and **1**B) shaped in accordance with embodiments of the technology. The superior portions **208** can have generally similar features as the superior portions **108** shown in FIGS. **1**A and **1**B. For example, each superior portion **208** includes an outer wall **212** and a septal wall **214**. The outer wall **212** is generally semi-circular, but can otherwise be configured according to the shape, geometry, and/or morphology of an arterial wall. The septal wall **214** can be shaped to mate with a complementary septal wall **214** of another endograft device. More specifically, in the embodiment illustrated in FIG. **2**A, the superior portion **208** includes a convexly curved, substantially semi-circular outer wall **212** and a substantially flat septal wall **214**. Thus, the superior portion **208** forms a "D" shape and can be part of a system (e.g., the system **100** shown in FIGS. **1**A and **1**B) including a corresponding D-shaped superior portion of a mating endograft device.

[0046] In other embodiments, both the outer wall **212** and the septal wall **214** can be convexly curved such that the superior portion **208** forms a complex ellipsoid with at least two distinct radii. FIG. **2**B, for example, shows the superior portion **208** can include a convexly curved outer wall **212** that has a first radius R**1** and a convexly curved septal wall **214** that has a second radius R**2** greater than the first radius R**1**. In the embodiment illustrated in FIG. **2**B, the second radius R**2** is substantially greater than the first radius R**1** such that the superior portion **208** has a substantially D-like shape.

[0047] Similarly, the superior portion **208** shown in FIG. **2**C includes the convexly curved outer wall **212** that has the first radius of curvature R**1** and the convexly curved septal wall **214** that has the second radius of curvature R**2** greater than the first radius R**1**. As shown in FIG. **2**C, the superior portion **208** can further include convexly curved corner sections **222** (identified individually as a first corner section **222***a* and a second corner section **222***b*). The first corner section **222***a* can have a third radius R**3**, and the second corner section **222***b* can have a fourth radius R**4** distinct from or equivalent to the third radius R**3**. In the embodiment shown in FIG. **2**C, the third and fourth radii R**3** and R**4** are substantially smaller than the first and second radii R**1** and R**2** such that the superior portion **208** forms another substantially D-like shape. In other

embodiments, the superior portion **208** can include greater or smaller radii, more or less curved portions, and/or can have another shape suitable for mating and at least substantially sealing two endograft devices together within a blood vessel.

[0048] FIGS. **2**D and **2**E are cross-sectional top views of the superior portion **208** of FIG. **2**B being mated with a complementary superior portion **208** to form a sealed septum **220** in accordance with an embodiment of the technology. More specifically, FIG. **2**D shows the superior portions **208** being pressed toward one another by a force F. The force F can derive from the self-expansion of the superior portions **208** within the confined space of an aorta. As shown in FIG. **2**D, the force F can cause the superior portions **208** to contact one another near the center of their respective convexly curved septal walls **214** and flatten the septal walls **214**. The apposition of the septal walls **214** can generate an outward force generally tangential to the septal walls **214** that can cause a slight outward bowing B near the interface of the outer and septal walls **212** and **214**.

[0049] As shown in FIG. **2**E, the force F can continue to press the superior portions **208** against one another until the convexly curved septal walls **214** straighten to form the septum **220**. The initial convexities of the septal walls **214** can induce more pressure between the septal walls **214** than straight septal walls (e.g., FIG. **2**A) and promote an even distribution of the force along the septum **220** to enhance the seal. Additionally, the outward bowing B can enhance the seal at the edges of the septal walls **214**. The superior portions **208** shown in FIGS. **2**A and **2**C can be similarly joined to form the substantially straight septum **220**. For example, the superior portion **208** shown in FIG. **2**C can be pressed against a corresponding superior portion such that the relative forces between the superior portions **208** substantially straighten the septal walls **214** and corner sections **222** (e.g., approximately 60° to 90° between the outer and septal walls **112** and **114**) to form the septum **220**. In operation, the septum **220** can be at least substantially sealed to prevent fluids (e.g., blood) from flowing between the superior portions **208**.

[0050] 1.3 Select Embodiments of Transition Portions

[0051] FIGS. **3**A and **3**B are isometric views of transition portions **324** of endograft devices configured in accordance with embodiments of the technology. The transition portions **324** can promote laminar blood flow by gradually changing the size of the lumen **116** from the wider, superior portion **108** to the narrower, inferior portion **110**. Additionally, the transition portions **324** can be configured to reduce the downforce exerted on the endograft devices **102** as blood flows through the lumen **116**.

[0052] More specifically, FIG. **3**A is an isometric view of the endograft device **102** described above with reference to FIGS. **1**A and **1**B. The endograft device **102** includes the transition portion **324** positioned between the superior portion **108** and the inferior portion **110**. As shown in FIG. **3**A, the transition portion **324** can be tapered to gradually narrow the cross-section of the lumen **116** and thereby reduce disruptions to the blood flow. The transition portion **324** can have a length L related to the distance necessary to continue substantially laminar blood flow through the lumen **116**. For example, in some embodiments, the length L can be 4 cm. In other embodiments, the length L can differ due to the geometry of the endograft device **102**, the rheologic characteristics of the blood flow, and/or other relevant factors in decreasing turbulent blood flow. In other embodiments, the transition portion **324** can be sloped, stepped, and/or have another suitable shape that can decrease the cross-section of the lumen **116** from the superior portion **108** to the inferior portion **110** without inducing turbulent blood flow.

[0053] FIG. **3**B is an isometric view of an endograft device **302** in accordance with another embodiment of the technology. The endograft device **302** can include generally similar features as the endograft **102** shown in FIG. **3**A. However, the tapered transition portion **324** shown in FIG. **3**B has a more gradual taper and a much greater length L than the transition portion **324** shown in FIG. **3**A. As shown in FIG. **3**B, the tapered transition portion **324** extends from the superior portion **108** to the second end portion **118***b* such that the transition portion **324** defines the inferior portion **110** (not visible). Accordingly, the tapered transition portion **324** can steadily decrease the cross-section of the lumen **116** to facilitate laminar blood flow through the lumen **116**. The gradual taper of the transition portion **324** may, however, cause the endograft device **302** to migrate in the direction of blood flow more than the more aggressive taper of the transition portion **324** shown in FIG. **3**A. Accordingly, the length L and angle of the tapered transition portion **324** can be optimized to mitigate migration of the endograft device **302** without inducing undo turbulent blood flow. In other embodiments, the transition portion **324** can optimize the geometry of a different shape (e.g., stepped) to maintain laminar blood flow and mitigate migration of the endograft device **302**.

[0054] 2. Endograft System Components

[0055] 2.1 Integrated Frames

[0056] FIGS. **4**A and **4**B are side views of the integrated frame **104** described with reference to FIGS. **1**A and **1**B in an expanded configuration (FIG. **4**A) and a low-profile configuration (FIG. **4**B) in accordance with an embodiment of the technology. As discussed above, the frame **104** includes the superior portion **108**, the inferior portion **110**, and the exposed end portions **118**. In some embodiments, the smallest radius of the outer wall **112** of each superior portion **108** in the expanded configuration may not be less than 10 mm (i.e., the smallest diameter of the superior portions **108** of mated endograft devices **102** is more than 20 mm).

[0057] As shown FIGS. **4**A and **4**B, the frame **104** can be a braided structure made from one or more continuous, interwoven wires **426** that provide a continuous, integrated support longitudinally along the length of the frame **104**. For example, as shown in FIG. **4**A, the wire **426** is braided such that a first longitudinal segment L**1** of the frame **104** supports an adjacent second longitudinal segment L**2** of the frame **104**. Accordingly, each area of the frame **104** influences the radial expansion or contraction of an adjacent area of the frame. In some embodiments, the frame **104** is woven with one wire **426** that continuously crosses itself along the length of the frame **104**. The intersections of the wire **426** may not be welded or otherwise fixed together such that they remain unbound to increase the flexibility of the frame **104**. In other embodiments, the frame **104** includes a plurality of wires **426** that can be interwoven and/or concentrically layered to form the frame **104**. The frame **104**, for example, can include eight wires **426** in which several of the wires **426** can end at intermediate points along the length of the frame **104**. Such a staggered, multi-wire construction prevents the wire ends from weakening the frame **104** and/or from wearing on a subsequently attached cover (e.g., the cover **106** shown in FIGS. **1**A and **1**B). The number of wires **426** can also vary at different sections along the length of the frame **104**. For example, in one embodiment, the inferior portion **110**

includes fewer wires **426** than the superior portion **108** such that the density or pitch of the wires **426** does not increase at inferior portion **110** and the frame **104**. This enables the inferior portion **110** to have a small diameter in the constricted, low-profile configuration (FIG. **4**B).

[0058] As shown in FIG. **4**A, the wires **426** can form a loop **428** at one end portion **118** to reverse direction and continue weaving along the length of the frame **104** toward the opposite end portion **118**. The optimal number of loops **428** at each end portion **118** can be associated with the diameter of the wires **426**. Too few loops **428** can decrease the strength at the end portions **118** of the contracted frame **104** shown in FIG. **4**A. Too many loops **428** can increase the profile of the extended frame **424** shown in FIG. **4**B, and can also cause difficulty attaching the cover. A wire **426** with a diameter of 0.008 inch, for example, may have an optimal number of ten to twelve loops **428** (five to six at each end portion **118**), whereas a wire **426** with a diameter of 0.009 inch may have an optimal number of twelve to fourteen loops **428**. In other embodiments, the wires **426** can include more or less loops **428** to optimize characteristics of the frame **104**. Additionally, the degree of curvature of each of the loops **428** can impact the durability of the wires **426**. For example, tightly wound loops **428** with high degrees of curvature are subject to fatigue and failure at the end portions **118** because of the stress induced upon constriction. Therefore, in some embodiments, the degree of curvature of the loops **428** can be the least degree of curvature permissible for the optimal number of loops **428**.

[0059] In the expanded configuration shown in FIG. **4**A, the wires **426** can cross at a braid angle θ selected to mitigate kinking and provide adequate extension/constriction. Lower braid angles θ can reduce or eliminate kinking of the wires **426** when the frame **104** is flexed or bent. For example, a braid angle θ of less than 45° allows the frame **104** to bend with smaller radii of curvature without substantial reduction of its cross-sectional area along the length of the frame **104**. Therefore, a frame **104** with a braid angle θ of less than 45° can be flexed and bent within the anatomy (e.g., the aorta) without restricting blood flow through the frame **104**. Additionally, lower braid angles θ can increase the outward spring force (i.e., the inherent force within the frame **104** that self-expands the frame **104** to the expanded configuration) and hoop strength (i.e., the radial strength of the frame **104** that restricts kinking and maintains the expanded configuration) of the frame **104**. Therefore, braid angles θ of not more than 45° can also provide an advantageous increase in the strength and corresponding durability of the frame **104**.

[0060] Lower braid angles θ, however, can also adversely affect the extension and constriction of the frame **104** in the low-profile configuration shown in FIG. **4**B. For example, extension and constriction can be negatively impacted at braid angles θ of less than 30°. Therefore, in some embodiments, the frame **104** can include a braid angle θ between 30° and 45° that promotes kink resistance and frame strength, while also maintaining extension and constriction abilities necessary for the low-profile configuration. In other embodiments, the optimal braid angle θ can be higher or lower.

[0061] In some embodiments in accordance with the technology, the braid angle θ can vary along the length of the frame **104** to vary kink resistance, outward spring force, hoop strength, and extension properties at different portions of the frame **104**. For example, the braid angle θ can be higher at the superior portion **108** (e.g., 40°) such that the superior portion **108** can extend and constrict into the low-profile configuration, and the braid angle θ can be lower at the inferior portion **110** (e.g., 30°) to provide kink resistance where the frame **104** is most likely to bend (e.g., within the aneurysmal sac and toward the iliac arteries). The smaller braid angle θ at the inferior portion **110** may not adversely affect the profile of the frame **104** because the inferior portion **110** need not constrict as much as the superior portion **108** to reach the desired low-profile configuration. In other embodiments, the braid angle θ of the frame **104** may vary in another way.

[0062] The wires **426** can have a diameter sufficient to support the frame **104** while still providing substantial flexibility for the frame **104**. The diameter of the wires **426** can be selected to attain a desired cross-sectional dimension in the low-profile configuration, a desired outward spring force to self-expand to the expanded configuration, and a desired hoop strength to support the frame **104** in the expanded configuration. For example, in some embodiments, the wires **426** can have a diameter from approximately 0.007 inch to approximately 0.014 inch. In specific embodiments, the wires have a diameter from approximately 0.011 inch to 0.013 inch. In other embodiments, the wires **426** can have a smaller diameter, a greater diameter, and/or the diameter of the wires **426** can vary along the length of the frame **104**. For example, in one embodiment, the wires **426** can have a greater diameter at the superior portion **108** than at the inferior portion **110** such that the wires **426** of the superior portion **108** have a outward spring force and greater hoop strength where the first and second endograft devices mate (e.g., at the septal walls **114**) and the increased density of wires **426** at the inferior portion **110** does not negatively impact the flexibility of the frame **104**.

[0063] The frame **104** may be constructed from a variety of resilient metallic materials, polymeric materials (e.g., polyethylenes, polypropylenes, Nylons, PTFEs, and the like), and composites of materials. For example, the wires **426** can be made from biocompatible stainless steels, highly elastic metallic alloys, and biocompatible shape setting materials that exhibit shape memory properties. In some embodiments, for example, the wire **426** can be made from a shape setting alloy, such as Nitinol, that has a preferred or native configuration. For example, a Nitinol structure can be deformed or constrained into a secondary configuration, but upon release from the constraint, the structure returns toward its native configuration with high fidelity. Accordingly, a frame **104** made from Nitinol wires **426** can reliably self-expand from the low-profile configuration the expanded configuration (i.e., its native configuration).

[0064] For endovascular delivery of a device (e.g., the endograft devices **102** shown in FIGS. **1**A and **1**B), the frame **104** is extended to constrict the frame **104** into a low-profile configuration in which the frame **104** can be loaded into a delivery device. The braid angle θ of the wires **426** can facilitate significant extension of the frame **104** to produce a slender profile during delivery as described above, and yet the interwoven characteristic of the braid restricts over extension. This extension-constriction functionality of the frame **104** allows the frame **104** to have variable diameters (e.g., the diameter of the superior portion **108** compared to the diameter of the inferior portion **11**) using the same number of wires **426** on each portion of the frame **104** such that the frame **104** has a low introduction profile (e.g., diameter) along the length of the frame **104**. The frame **104** can also include an optimal

number of loops **428** at each end portion **118** such that the loops **428** do not increase the profile of the frame **104** upon full extension.

[0065] At a target site (e.g., above an aneurysm), the frame **104** self-expands to the expanded configuration shown in FIG. **4**A as it is removed from the delivery device. The braid angle θ can be adjusted to change the outward spring force and hoop, strength of the expanded frame **104** as explained above. In some circumstances, the endograft device may need to be repositioned after being partially deployed. The frame **104** is well suited for such repositioning because the loops **428** and the continuous, interwoven wires **426** can simplify recapture of the frame **104** and allow for constriction after expansion to correctly reposition the endograft device. Additionally, portions of the frame **104** can remain exposed (e.g., the end portions **118**) to encourage cell ingrowth for securely anchoring the frame **104** to the arterial walls. Moreover, as described in more detail below, the interwoven wires **426** of the braided frame **104** can provide a continuous longitudinal support along the length of the frame **104** such that the frame **104** can be staggered and free end portions can support themselves. The frame **104** can also facilitate attachment to other endograft devices. For example, the frame **104** can interlace with another interwoven wire **426** of a supra-renal endograft.

[0066] Once deployed across the aneurysm, the frame **104** can also accommodate disparate anatomies and morphologies. In several patients, the aneurysmal sac extends at an angle with respect to the neck of the aneurysm. Because the frame **104** can have a braid angle θ that prevents kinking, the frame **104** can bend and flex without kinking to accommodate angulated aneurysmal sacs without restricting blood flow. Additionally, the unbound, woven wires **426** give the frame **104** a radial elasticity such that the frame **104** mimics the changes in the shape and morphology of the aorta without hindering the interface or seal between the endograft device and the vessel wall. For example, the frame **404** can constrict and expand to maintain the seal when pressure and other conditions alter the vasculature of the aorta. Moreover, the woven wires **426** inherently generate a spring force that biases the frame **104** toward a substantially straight trajectory within an aneurysmal sac and thereby limits migration of the endograft device.

[0067] In addition, the constant outward spring force and hoop strength of the braided frame **104** can be adjusted by changing the braid angle θ and/or the diameter of the wires **426**. This allows the formation of large diameter frames **104** without a significant change in the low-profile cross-sectional dimensions. Additionally, this feature allows the frames **104** to contract to a much smaller introduction profiles (e.g., diameters) compared to standard Z-frames or M-frames because the standard Z-frames and M-frames tend to require more wire and therefore larger introduction profiles to maintain a constant outward spring force and hoop strength.

[0068] 2.2 Covers

[0069] FIGS. **5**A-C are views of a cover being extended from an expanded configuration (FIG. **5**A) to a low-profile configuration (FIG. **5**C) in accordance with embodiments of the technology. More specifically, FIG. **5**A is a side view of the cover **106** described above with reference to FIGS. **1**A and **1**B in the expanded configuration. The cover **106** can include a plurality of circumferential ribs **530** such that the cover **106** has an undulating profile. As shown in FIG. **5**A, the individual ribs **530** can have a substantially triangular shape with an apex **533**. In other embodiments, the individual ribs **530** have rounded edges, rectangular edges, and/or other suitable textures that can extend and contract.

[0070] The ribs **530** of one cover can mate with opposing ribs **530** of an opposing cover and interface with vessel walls to enhance the seal and fixation between endograft devices in an endograft system (e.g., the endograft devices **102** of the endograft system **100** shown in FIGS. **1**A and **1**B) and between the endograft devices and the arterial walls. For example, the apices **533** of the ribs **530** at the septal wall **114** of the superior portion **108** of one endograft device can interface or mate with the troughs of the corresponding ribs **530** on a cover of an opposing endograft device. Additionally, the ribs **530** at the outer wall **112** can contact the arterial walls in a manner that at least substantially seals them together. The ribs **530** can also allow the cover **106** to flex and bend without wrinkling in situ. In some embodiments, the ribs **530** can be at only selected portions of the cover **106** (e.g., the septal wall **114**). In other embodiments, the ribs **530** can have different shapes and/or geometries on different portions of the cover **106**. For example, the apices **533** of the ribs **530** can have a first height on the superior portion **108** to enhance sealing forces between the endograft devices and a second height less than the first height at the inferior portion **110** to allow the cover **106** to freely flex and bend to accommodate the anatomy.

[0071] The ribs **530** change with the expansion and contraction of the cover **106**. As shown in FIG. **5**A, the apices **533** of the ribs **530** protrude to the maximal extent in the expanded configuration. Referring to FIG. **5**B, as the cover **106** extends, the ribs **530** also extend and constrict. When the cover **106** is fully extended in the low-profile configuration shown FIG. **5**C, the ribs **530** are completely elongated and constricted. In some embodiments, the size of each rib **530** can be predetermined to ensure the ribs **530** are completely flattened in the low-profile configuration and project radially outwardly to interface with adjacent surfaces in the expanded configuration. Accordingly, the ribs **530** do not limit the mobility of the endograft device as it is delivered to the aorta in the low-profile configuration.

[0072] Additionally, as shown in FIGS. **5**A-C, the cover **106** can include zigzagged edges at a superior terminus **531***a* and an inferior terminus **531***b* of the cover **106**. The zigzagged termini **531** can facilitate substantially seamless attachment between the cover **106** and an integrated frame (e.g., the frame **104** shown in FIGS. **4**A and **4**B). For example, in some embodiments, the zigzagged termini **531** can correspond to the braid angle θ of interwoven wires. The zigzagged termini **531** generally prevent the cover **106** from wrinkling or bunching at first and second end portions (e.g., the first and second end portions **118***a* and **118***b* shown in FIGS. **4**A and **4**B) when the cover **106** and the frame are constricted. In other embodiments, the superior and inferior termini **531***a* and **531***b* can be scalloped, straight, and/or have another suitable shape that facilitates attachment and/or limits wrinkling.

[0073] The cover **106** can be made from a substantially impermeable, biocompatible, and flexible material. For example, the cover **106** can be made from synthetic polymers, polyurethanes, silicone materials, polyurethane/silicone combinations, rubber materials, woven and non-woven fabrics such as Dacron®, fluoropolymer compositions such as a polytetrafluoroethylene (PTFE) materials, expanded PTFE materials (ePTFE) such as TEFLON®, GORE-TEX®, SOFTFORM®, IMPRA®, and/or other suitable materials. Additionally, in some embodiments, the cover **106** can be

made from a material that is sufficiently porous to permit ingrowth of endothelial cells. Such a porous material can provide more secure anchorages of endograft devices and potentially reduce flow resistance, sheer forces, and leakage of blood around the endograft devices.

[0074] In some embodiments in accordance with the technology, the cover **106** may also include drug-eluting coatings or implants. For example, the cover **106** can be coated and/or imbedded with a slow-releasing drug that can block cell proliferation, promote reendothelialization of the aneurysm, and/or otherwise medicate the aneurysmal region. Suitable drugs can include calcium, proteins, mast cell inhibitors, and/or other suitable medicines that encourage beneficial changes at the aneurysmal region.

[0075] In accordance with other embodiments of the technology, the cover **106** can be eliminated in favor of one or more layers of a coating material (shown and described in more detail with reference to FIGS. **17**A-E). The coating layer can be made from a biocompatible synthetic polymer, such as PTFE. The coating layer can be placed on the interior of an integrated frame (e.g., the frame **104** shown in FIGS. **4**A and **4**B), the exterior of the frame, and/or interwoven throughout the frame. Like the cover **106**, the coating layers can encase the frame to form a lumen (e.g., the lumen **116** shown in FIGS. **1**A and **1**B). Additionally, the coating can have a selected porosity that encourages tissue ingrowth.

[0076] 2.3 Integrated Frame and Cover

[0077] FIGS. **6**A and **6**B are cross-sectional views of the endograft device **102** of FIGS. **1**A and **1**B in a low-profile configuration and an expanded configuration, respectively, in accordance with embodiments of the technology. As shown in FIGS. **6**A and **6**B, the cover **106** can be attached to the exterior of the frame **104** at one or more attachment areas **632** (identified individually as a first attachment area **632***a* and a second attachment area **632***b*). The attachment areas **632** can have sutures, adhesives, welds, and/or other suitable fasteners that discretely hold the cover **106** to the frame **104** at the attachment areas **632**.

[0078] In the embodiment shown in FIGS. **6**A and **6**B, the endograft device **102** has attachment areas **632** at only the superior and inferior termini **531***a* and **531***b* of the cover **106** such that the remainder of the cover **106** between the attachment areas **632** is not attached directly the frame **104**. As a result, the frame **104** and the cover **106** can fully extend and constrict as shown in FIG. **6**A without interfering with one another. For example, in the low-profile configuration shown in FIG. **6**A, the frame **104** does not directly pull the central portion of the cover **106** downward and longitudinally with the frame **104** such that the ribs **530** can stretch uniformly along the length of the cover **106** to accommodate full extension of the frame **104**. Similarly, the intermediate portions of the cover **106** do not hinder the extension or constriction of the frame **104**. Fewer attachments areas **632** can also limit the potential for fatigue and undesirable porosity that may arise at the attachment areas **632**, such as from needle pricks and other fastening mechanisms that puncture the cover **106**.

[0079] As shown in FIG. **6**B, the cover **106** can substantially conform to the shape of the frame **104** when they are in the expanded configuration. Proper alignment between the cover **106** and the frame **104** prevents the cover **106** from adversely affecting constriction and expansion. For example, alignment between the cover **106** and the frame **104** at the superior and transition portions **108** and **324**, respectively, ensures the frame **104** can expand properly and generate the force necessary to mate with a superior portion of an opposing endograft device. Additionally, in some embodiments, the cover **106** is sized to restrict the expansion and corresponding contraction of the frame **104**.

[0080] Attaching the cover **106** to the exterior of the frame **104** as shown in FIGS. **6**A and **6**B can provide a plurality of benefits for the endograft device **102**. For example, unlike endograft devices with internal covers that must fold within a frame during delivery, the exterior cover **106** does not inhibit constriction of the frame **104** (e.g., FIG. **6**A). In the expanded configuration, the exterior the cover **106** does not bunch or wrinkle within the frame **104**, and thus does not cause thrombotic problems within the lumen **116**. Additionally, unlike more rigid Z-stents, the flexibility of the frame **104** can prevent abrasive rubbing and deterioration of the cover **106** in the expanded configuration (e.g., FIG. **6**B). The exterior attachment of the cover **106** can also prevent over expansion of the frame **104**.

[0081] 2.4 Alignment Aids

[0082] FIGS. **7**A and **7**B are isometric views of endograft devices **702** in accordance with additional embodiments of the technology. The endograft devices **702** can have generally similar features as the endograft devices **102** shown in FIGS. **1**A and **1**B. Additionally, the endograft devices **702** can include alignment aids **734** that are visible under imaging systems (e.g., X-rays) to facilitate accurate positioning and subsequent monitoring of the endograft devices **702** in the vasculature.

[0083] FIG. **7**A is a partial cut-away isometric view of the endograft device **7-2** showing an alignment aid **734** in accordance with an embodiment of the technology. As shown in FIG. **7**A, the alignment aid **734** can extend diagonally along the septal wall **114** of the frame **104** to indicate the position of the septal wall **114** relative to the endograft device **702**. The alignment aid **734** can thus provide an indication of the rotational orientation and axial location of the endograft device **702** such that during deployment opposing septal walls **114** can be properly aligned and mated with one another. Additionally, as shown in the embodiment in FIG. **7**A, the alignment aid **734** can terminate at the superior terminus **531***a* of the cover **106** to indicate where the first end portion **118***a* begins. Thus, the alignment aid **734** provides a definitive indicator to ensure that the cover **106** does not block transverse flow (e.g., from the aorta to the renal arteries). In other embodiments, the alignment aids **734** may be positioned elsewhere along the endograft device **702** to provide spatial location and orientation that can aid delivery and deployment of the endograft device **702**.

[0084] The alignment aid **734** can be made from radiopaque and/or fluoroscopic materials, such as tantalum, platinum, gold, and/or other materials that are visible under an imaging system (e.g., X-rays). For example, as shown in FIG. **7**A, the alignment aid **734** is made from a radiopaque wire (e.g., tantalum) wound around a segment of the frame **104**. In another embodiment, a radiopaque composition is applied to the frame **104** and/or incorporated in the septal walls **114** of the cover **106**.

[0085] FIG. **7**B shows the first and second endograft devices **702** mated together using the alignment aids **734** in accordance with an embodiment of the technology. As shown in FIG. **7**B, the alignment aids **734** on the first and second endograft devices **702***a* and **702***b* are symmetrical such that when the endograft devices **702** are correctly oriented and the septal walls **114** oppose one another, the alignment aids **734**

can intersect to form an "X" indicator. In other embodiments, the intersection of the alignment aids **734** forms other characters, numbers, and/or symbols that indicate the rotational orientation and longitudinal location of the endograft devices **702**. In further embodiments, the alignment aids **734** can be applied to different portions of the septal wall (e.g., the cover **102**) and/or the outer wall **112**. In still further embodiments, the endograft devices **702** include a plurality of alignment aids **734** to distinguish different portions of the endograft devices **702** and further aid rotational and/or other orientation. For example, in some embodiments, the inferior portions **110** include alignment aids **734** that differentiate the inferior portions **110** of the first and second endograft devices **702**.

[0086] 2.5 Anchors

[0087] FIGS. **8**A and **8**B are isometric views of endograft devices **802** configured in accordance with additional embodiments of the technology. The endograft devices **802** can include generally similar features as the endograft devices **102** shown in FIGS. **1**A and **1**B. Additionally, the endograft devices **802** can include one or more anchors **836** that project outwardly from the frame **104** and/or cover **106** to engage the interior surfaces of arterial walls. The anchors **836** can be barbs, hooks, and or other shapes that can penetrate into the arterial walls. For example, as shown in FIG. **8**A, the anchors **836** can be "V" shaped projections. In some embodiments, the anchors **836** eventually become embedded in cell growth on the interior surface of the arterial wall. In operation, the anchors **836** resist migration of the endograft devices **802** within the artery and reduce the likelihood of endoleaks between the outer wall **112** and the arterial wall.

[0088] In an embodiment shown in FIGS. **8**A and **8**B, the anchors **836** project from the outer walls **112** to secure the superior portions **108** to the aorta. In other embodiments, additional anchors **836** can project from the second end portions **118***b* to secure the inferior portions **110** to the iliac arteries. The anchors **836** can also protrude from the septal walls **114**, extend through the lumen **116**, and project outward beyond the outer wall **112** to enhance the strength of the engagement. The anchors generally project inferiorly such that downward forces applied to the endograft devices **802** (e.g., blood flow) drive the anchors **836** further into the arterial walls.

[0089] In one embodiment in accordance with the technology, the anchors **836** are separate elements that are attached to the frame **104**. For example, in the embodiment shown in FIG. **8**A, the anchors **836** are small barbs or wires that are fastened to the frame **104** by winding another wire (e.g., a Nitinol wire) around the anchors **836** and the adjacent wire **426** of the braid. In other embodiments, the anchors **326** are integrally formed with the wire **426** used in the braid of the frame **104**. For example, as shown in FIG. **8**B, the anchors **836** are woven into the outer wall **112** of the frame **104**. The interwoven anchors **836** can be deployed (i.e., project outwardly) when the frame **104** expands and can retract when the frame **104** constricts. Accordingly, the interwoven anchors **836** do not inhibit movement of the endograft device **802** during delivery in the low-profile configuration. In other embodiments, the anchors **836** can be attached to a different portion of the endograft device **802** (e.g., the cover **106**).

[0090] The anchors **836** can be made from resilient metallic materials, polymeric materials (e.g., polyethylenes, polypropylenes, Nylons, PTFEs), and/or other suitable materials that can anchor the endograft devices **802** to arterial walls. For example, the interwoven anchors **836** shown in FIG. **8**B can be made from Nitinol wire **426** that comprises the frame **104**.

[0091] 3. Methods of Implementation and Assembled Endograft Systems

[0092] Described below are methods of deploying and assembling modular endograft systems across an aneurysm in accordance with embodiments of the technology. The associated Figures (i.e., FIGS. **9**A, **9**B, **11**-**13**C and **15**A-**16**) include schematic representations of an abdominal portion of an aorta. More specifically, FIG. **9**A shows an aneurysm **50** located along an infrarenal portion of the aorta **52**, which is the most common site of an AAA. A right or first renal artery **54***a* and a left or second renal artery **54***b* stem from the aorta **52**. The region of the aorta **52** superior to the aneurysm **50** and inferior to the renal arteries **54** is the aortic neck **60**. The distal end portion of the aorta **52** bifurcates into common iliac arteries **56** (identified individually as a first iliac artery **56***a* and a second iliac artery **56***b*), and the internal iliac arteries **58** (identified individually as a first internal iliac artery **58***a* and a second internal iliac artery **58***b*) branch from the common iliac arteries **56**. Other arteries and structures proximate to the abdominal portion of the aorta **52** have been removed for clarity.

[0093] 3.1 Modular Endograft Systems

[0094] FIGS. **9**A and **9**B are schematic views of the two-part modular endograft system **100** described above being deployed across the aneurysm **50** in accordance with an embodiment of the technology. FIG. **9**A shows a delivery system **40** for implanting the first and second endograft devices **102***a* and **102***b*. The delivery system can include a first catheter **42***a*, a first guidewire **44***a* associated with the first catheter **42***a*, a second catheter **42***b*, and a second guidewire **44***b* associated with the second catheter **42***b*. Each endograft device **102** (FIG. **9**B) can be extended to the low-profile configuration and loaded into the corresponding catheter **42**. Because the endograft devices **102** are delivered separately, the sizes of the catheters **42** are not constrained by the system **100** as a whole. In some embodiments, for example, the low-profile configurations of each endograft device **102** can fit within a 12 F catheter. In other embodiments, the low-profile configuration of the endograft devices **102** can fit within differently sized catheters **42**.

[0095] During deployment, the first catheter **42***a* and the first guidewire **44***a* are inserted percutaneously into a blood vessel (e.g., a femoral artery; not shown). With the aid of imaging systems, the first guidewire **44***a* is endoluminally navigated through the vasculature, up the first iliac artery **56***a*, and to a location superior to a target site T above the aneurysm **50**. The first catheter **42***a* is then passed through the vasculature along the first guidewire **44***a* to the target site T. Using a generally similar method, the second guidewire **44***b* and the second catheter **42***b* are delivered through the second iliac artery **56***b* to the target site T. The first and second endograft devices **102***a* and **102***b* can be delivered simultaneously or in succession.

[0096] The endograft devices **102** can be urged out of the distal ends of the catheters **42** at the target site T by withdrawing the catheters **42** proximally while holding the endograft devices **102** in place using pushers or other suitable endovascular instruments. Alternatively, the endograft devices **102** can be pushed distally while holding the catheters **42** in place. Upon release, the endograft devices **102** self-expand to the expanded configuration shown in FIG. **9**B. The guidewires **44**

generally remain in place to facilitate adjusting the endograft devices **102**. This eliminates the need to cannulate either of the endograft devices **102**.

[0097] Each endograft device **102** can be positioned at its desired location independently of the other endograft device **102** while the endograft devices **102** are in, or at least partially within, the catheters **42**. For example, in the embodiment illustrated in FIG. **9**B, the superior portions **108** contact the aortic neck **60** at the same level, and the inferior portions **110** extend through the aneurysm **50** to their respective iliac arteries **56**. More specifically, the inherent hoop force of the frame **104** caused by the constant outward spring force of the braid at least substantially seals (a) the covers **106** at the outer walls **112** against the aortic neck **60** and (b) the septal walls **114** to each other to form the septum **120**. The inferior portions **110** extend through the aneurysm **50** and can bend to enter the iliac arteries **56**. The proximal portion of the inferior portions **110** contact the iliac arteries **56** and can form a seal therebetween. The flexibility of the frame **104** prevents the endograft devices **102** from kinking at the bend and restricting blood flow. Additionally, as shown in FIG. **9**B, the spring force within the frame **104** biases the inferior portions **110** to extend in a substantially straight trajectory through the aneurysm **50**. This inhibits migration of the inferior portions **110** to a side of the aneurysm **50** that could break the contact and/or seal at the aortic neck **60**. As described in more detail below, in other embodiments the endograft devices **102** can be positioned independently at different elevations along the aortic neck **60**.

[0098] As further shown in FIG. **9**B, the endograft system **100** can include extension units **937** (identified individually as a first extension unit **937***a* and a second extension unit **937***b*) projecting distally from the superior termini **531** of the covers **106**. The extension units **937** can include an extension frame **904** (not visible) and an extension cover **906** at least generally similar to the frame **104** and the cover **106** of the endograft devices **102** described above. The extension units **937** can have a substantially similar shape as the superior portions **108** of the endograft devices (e.g., a D-like shape) such that the extension units **937** can mate with the interior of at least a part of the superior portions **108**. For example, as shown in FIG. **9**B, the extension covers **906** can be positioned inferior to the renal arteries **54** within the frame **104** such that the extension covers **906** can interface with the aortic neck **60** and mate with one another to extend the septum **120** distally. Therefore, the extension units **937** can increase the fixation area and the sealing area of the endograft devices **102** when the superior termini **531** of the covers **106** of the endograft devices **102** are offset from the entrances of the renal arteries **54**. For example, in some embodiments, the extension units **937** add approximately one inch of fixation structure and sealing area to the endograft devices **102**. In other embodiments, the inferior portions **110** can also include extension units **937** that can affix and at least substantially seal to the iliac arteries **56**.

[0099] During deployment, the extension units **937** can be added to the system **100** after the first and second endograft devices **102** are positioned within the aortic neck **60**. With the aid of the delivery system **40**, the extension units **937** can advance along the guidewires **44** and be deployed from the catheters **42** at desired positions within the first and second frames **104** just inferior of the renal arteries. Upon deployment, the extension units **937** can self-expand via an inherent spring force in the extension frame **904** to an expanded configuration to contact and at least substantially seal with the interior of the superior portions **108** of the endograft devices **102**. As shown in FIG. **9**B, the extension cover **906** can interface with the first end portions **118***a* of the frames **104** to strengthen the seal therebetween. In other embodiments, the extension units **937** can connect and seal to the endograft devices **102** using other suitable attachment methods. The extension units **937** can be positioned independently such that they accommodate anatomical variations (e.g. staggered renal arteries). For example, a superior terminus of the first extension unit **937***a* can be longitudinally offset from a superior terminus of the second extension units **937***b*. Similarly, the inferior portions **110** can include extension units **937** that increase the sealing area with the iliac arteries **56**.

[0100] In some embodiments, alignment aids, such as the alignment aids **734** described with reference to FIGS. **7**A and **7**B, are used to rotationally orient the endograft devices **102** and align the septal walls **114** during delivery. Additionally, to prevent migration and/or projection of the system while in situ, anchors, such as the anchors **836** described above with reference to FIGS. **8**A and **8**B, can be deployed from the outer walls **112** to engage the arterial walls of the aortic neck **60** and/or from the second end portions **118***b* to engage the arterial walls of the iliac arteries **56**.

[0101] FIGS. **10**A-**11** show additional embodiments of implementing endograft systems (e.g., the system **100**) in which the superior portions **108** are longitudinally offset from each other. For example, in some embodiments, the superior portions **108** are longitudinally offset by at least 5 mm. The features of the systems below allow one or both of the superior portions **108** to be placed over transverse arteries to increase the available fixation structure and sealing area for the endograft devices **102** without inhibiting blood flow.

[0102] FIG. **10**A is an isometric view of the modular endograft system **100** in which the endograft devices **102** are staggered such that the superior portion **108** of the first endograft device **102***a* is above the superior portion **108** of the second endograft device **102***b*. The first end portion **118***a* of the second endograft device **102***b* can prevent the unsupported free first end portion **118***a* of the first endograft device **102***a* from splaying outward into the blood flow in a manner that induces undo turbulence. Moreover, the interplay between the woven wires **426** of the frame **104** of the first endograft device **102***a* restricts the outward movement of the first end portion **118***a* of the first endograft device **102***a* and provides substantially continuous support along the length of the frame **104** such the free first end portion **118***a* retains substantially the same shape as if it were supported. These features maintain the generally straight or convex shape of the unsupported septal region of the first portion **118***a* of the first endograft device **102***a*. Using shape-setting Nitinol wire **426** in the frame **104** can further facilitate maintaining the shape of the unsupported portion of the frame **104**.

[0103] Compared to conventional devices that have a common height across the diameter of a vessel (e.g., the aorta), the staggered configuration shown in FIG. **10**A allows one or both of the first end portions **118***a* to extend over the entrance of the renal arteries to increase the available structure for fixing the endograft devices **102** to the vessel wall. The staggered configuration also increases the sealing area of the superiorly positioned first endograft device **102***a* for anatomies having a short aortic neck (e.g., less than 2 cm). Similarly, the second end portions **118***b* can extend over the entrances of the internal iliac arteries to ensure the inferior portions **110** each have an adequate structure for fixing and at

least substantially sealing the inferior portions **110** to the iliac arteries. To the extent migration occurs, the additional sealing area between the endograft devices **102** and the vessel walls will reduce the potential for leakage at the aortic neck.

[0104] FIG. **10**B is an isometric view of a modular endograft system **1000** configured in accordance with an additional embodiment of the technology. The system **1000** can have a first endograft device **1002***a* and a second endograft device **1002***b* that are generally similar to the endograft devices **102** described above. The covers **106** of the endograft devices **1002** in FIG. **10**B, however, extend to the distal ends of the superior portions **108**. Additionally, the endograft devices **1002** further include fenestrations **1038** on the outer walls **112** of the superior portions **108**.

[0105] The fenestrations **1038** can be openings through the cover **106** that expose the frame **104** and provide a channel through which blood can flow to and from transverse arteries. For example, the endograft devices **1002** can be positioned independently and staggered such that the fenestration **1038** of each endograft device **1002** is aligned with one of the left or right renal arteries. The fenestrations **1038** accordingly increase the available sealing area between the outer walls **112** and the arterial walls because the superior portions **108** can be positioned independently over the renal arteries such that one endograft device **1002** does not need to be limited to the elevation of the inferior renal artery. This provides optimal placement for each endograft device **1002** within the vasculature without requiring customized devices. In other embodiments in accordance with the technology, the endograft devices **1002** can include additional fenestrations **1038** to increase the available sealing area without restricting blood flow. For example, the inferior portions **110** can include fenestrations **1038** that allow the inferior portions **110** to extend over the entrance of the internal iliac arteries.

[0106] FIG. **11**A is a schematic view of the modular endograft system **100** deployed across an aneurysm such that the superior portions **108** of the endograft devices **102** are staggered to accommodate for anatomical variations in the vasculature in a manner that takes advantage of the available structure for fixing the endograft devices **102** to arterial walls and the available sealing area in the aortic neck **60**. In the embodiment shown in FIG. **11**A, for example, the left renal artery **54***b* is inferior the right renal artery **54***a*. The first endograft device **102***a* can, therefore, also be positioned higher in the aorta **52** to utilize the available fixation and sealing areas on the ipsilateral side of the aortic neck **60** without having to be concerned about blocking the entrance of the left renal artery **54***b*. The first end portion **118***a* of the second endograft device **102***b* can be positioned over the left renal artery **54***b* without inhibiting blood flow to lengthen the structure for fixing the second endograft device **102***b* to the arterial wall and mating the septal walls **114** together. The longer fixation and sealing areas along the outer wall **112** of the first endograft device **102***a* and the longer mating and sealing areas between the septal walls **114** can strengthen the seals of the system **100** as a whole to reduce the likelihood of endoleaks. Additionally, as shown in FIG. **11**A, the system **100** can be staggered to accommodate an anatomy with less fixation and sealing area in one of the iliac arteries **56**.

[0107] FIG. **11**B is a schematic view of the modular endograft system **1000** of FIG. **10**B deployed across the aneurysm **60**. Similar to the configuration of the system **100** shown in FIG. **11**A, the endograft devices **1002** are staggered to accommodate for anatomical variations in the vasculature in a manner that takes advantage of the available anatomical structure for fixing and sealing the outer walls **112** of the endograft devices **102** to the arterial walls in the aortic neck **60**. As shown in FIG. **11**B, for example, the first endograft device **1002***a* can be positioned superior to the second endograft device **1002***b* in the aortic neck **60** to utilize the available fixation and sealing area on the ipsilateral side of the aortic neck **60**. The fenestrations **1038** can be placed independently at the entrance of each renal artery **54** to increase the available fixation and sealing area in the aortic neck **60** and accommodate asymmetrical anatomies. Additionally, as further shown in FIG. **11**B, the endograft devices can include fenestrations **1038** at the inferior portions **110** that can be placed independently at the entrance of each internal iliac artery **58** to accommodate an anatomy with less sealing area in the iliac arteries **56**. In other embodiments, the endograft devices **102** can include fenestrations **1038** to accommodate other anatomical variations.

[0108] FIG. **12** is a schematic view of the modular endograft system of FIGS. **9**A and **9**B deployed across an angulated aneurysm in accordance with an additional embodiment of the technology. The system **100** can accommodate this anatomical abnormality because the endograft devices **102** are flexible. More specifically, the interwoven wires **426** of the frame **104** are sufficiently flexibility to bend without kinking. Thus, the bent endograft devices **102** can maintain unrestricted flow through the lumens **116**. Accordingly, the system **100** can accommodate other anatomical variations that may require the endograft devices **102** to flex or bend without disturbing blood flow.

[0109] FIGS. **13**A-C are schematic views of a four-part modular endograft **1300** system ("system **1300**") being deployed across the aneurysm **50** in accordance with an embodiment of the technology. The system **1300** can include generally similar features as the system **100** described with reference to FIGS. **9**A and **9**B. However, as shown in FIG. **13**B, the inferior portions **110** of the endograft devices **102** terminate within the aneurysm **50**. Therefore, as shown in FIG. **13**C, the system **1300** further includes separate limbs **1362** (identified individually as a first limb **1362***a* and a second limb **1362***b*) that contact and substantially seal with corresponding inferior portions **110** and extend into corresponding iliac arteries **56**. The limbs **1362** can be generally similar to the inferior portions **110**. For example, the limbs **1362** can include an integrated frame **1304** and a cover **1306** generally similar to the frame **104** and the cover **106** described above with reference to FIGS. **1**A-**6**B. As shown in FIG. **13**C, the limbs **1362** self-expand within the interior portions **110** to the expanded configuration and thereby the superior portions of the limbs **1362** at least substantially seals to the proximal section of the inferior portions **110**. The length of the limbs **1362** within the inferior portions **110** can be adjusted to increase the available structure for fixing and sealing the limbs **1362** to the endograft devices **102**. Additionally, in some embodiments, the covers **1306** of the limbs **1362** can include ribs, such as the ribs **530** described above with reference to FIGS. **5**A-C, that interface with the interior of the frames **104** and the covers **106** at the inferior portions **110** to connect and at least substantially seal the limbs **1362** to the inferior portions **110**. In other embodiments, the limbs **1362** can connect and at least substantially seal to the exteriors of the inferior portions **110** using anchors (e.g., the anchors **836** described with reference to FIGS. **8**A and **8**B), self-constricting forces, and/or other suitable attachment and

sealing methods. The limbs **1362** extend the lumens **116** of the endograft devices **102** to the iliac arteries **56** such that blood can flow through the system **1300** to bypass the aneurysm **50**.

[0110] Referring to FIG. **13**A, the delivery system **40** is shown within the abdominal portion of the aorta **52** before deploying the endograft system **1300**. The insertion of the delivery system **40** can be generally similar as described above with reference to FIG. **9**A. However, as shown in FIG. **13**A, the first and second guidewires **44***a* and **44***b* can cross after they enter the aneurysm **50** such that each catheter **42** extends from its respective iliac artery **54** to the contralateral side of the aorta **52**. For example, the first catheter **42***a* can be delivered from the first iliac artery **56***a* to the left side of the aorta **52** proximate to the left renal artery **54***b* (Arrow D_1), and the second catheter **42***b* can be delivered from the second iliac artery **56***b* to the right renal artery **54***a* (Arrow D_2). In other embodiments, such as in the deployment method described above with reference to FIGS. **9**A and **9**B, the guidewires **44** do not cross within the aneurysm **50**.

[0111] Referring to FIG. **13**B, after the first and second catheters **42***a* and **42***b* are positioned in the aortic neck **60**, they are pulled proximally to deploy the endograft devices **102** through the distal ends of the catheters **42**. The crossing catheters **42** and guidewires **44** deploy the endograft devices **102** on opposite sides of the aortic neck **60**.

[0112] As shown in FIG. **13**B, the inferior portions **110** of the endograft devices **102** terminate within the aneurysm **50** and form a "gate." In general, gates are considered undesirable because in conventional systems they must be cannulated to deliver and deploy limbs that extend the endograft devices into the iliac arteries **56**. However, as shown in FIG. **13**B, the guidewires **44** remain within the endograft devices **102** after they are deployed; this eliminates the need for time-consuming cannulation of the gates because the inferior portions **110** of the endograft devices **102** are in effect pre-cannulated. Such pre-cannulated gates allow the limbs **1362** to be delivered through the distal ends of the catheters **42** and connected to the inferior portions **110** much faster and more accurately than conventional systems.

[0113] FIG. **13**C shows the system **1300** after both limbs **1362** are connected to the endograft devices **102**. As shown in FIG. **13**C, the delivery system **40** can also be used to adjust the length of the limbs **1362** and the length of the fixation area between the limbs **1362** and the inferior portions **110** in the direction of the arrows. In the embodiment shown in FIG. **13**C, for example, the second limb **1362***b* extends further into the inferior portion **110** of the second endograft device **102***b* such that the second limb **1362***b* is effectively shorter than the first limb **1362***a*. The length of the limbs **1362** can be adjusted to accommodate disparate anatomies of the iliac arteries **56**, maximize the fixation and sealing areas of the limbs **1362**, and/or otherwise optimize the position of the limbs **1362**. This is possible because, at least in part, the inferior portions **110** of the endograft devices **102** can be relatively long to allow significant longitudinal leeway in positioning the limbs **1362** while still providing adequate surface area to at least substantially seal the limbs **1362** to the inferior portions **110**.

[0114] The four-part, two-wire system **1300** can easily accommodate anatomical variations without requiring customized components. For example, the superior portions **108** can be staggered to maximize the mating and sealing area of each outer wall **112** with the aortic walls. Additionally, each limb **1362** can be selected from a relatively small number of different lengths to extend a desired length within the iliac arteries **56** that both adequately connects and substantially seals the limbs **1362** to the arterial walls and does not block transverse arterial flow. The limbs **1362** can also be adjusted independently relative to the inferior portions **110** to increase the available structure for fixing and sealing the limbs **1362** and the inferior portions **110** together, and to shorten or lengthen the limbs **1362** within the iliac arteries **56**. Additionally, the braided structure of the frames **104** can decrease infolding of the covers **106** such that the lengths of the frame **104** can be selected from standardized cross-sectional dimensions. Thus, the four-part system **1300** can be highly customizable, but yet comprise standardized components.

[0115] 3.2 Modular Endograft System with Aortic Cuff

[0116] FIGS. **14**A and **14**B are isometric views of a modular endograft system **1400** ("system **1400**" shown in FIG. **14**B) configured in accordance with embodiments of the technology. More specifically, FIG. **14**A is an isometric view of an aortic cuff **1464** for use with the endograft devices **102** (FIG. **14**B). The aortic cuff **1464** can include a sleeve **1466** and a cuff frame **1468**. As shown in FIG. **14**A, the sleeve **1466** and the cuff frame **1468** can be separate components. In other embodiments, the sleeve **1466** and the cuff frame **1468** can be formed integrally. The aortic cuff **1464** can expand from a low-profile configuration having a first cross-section to an expanded configuration (e.g., FIG. **14**B) having a second cross-section larger than the first cross-section. The low-profile configuration can be used during delivery of the aortic cuff **1464** from which the cuff-device **1464** can self-expand to the expanded configuration in situ. The aortic cuff **1464** can be configured to interface and substantially seal with an infrarenal portion of the aorta superior to an aneurysm.

[0117] The sleeve **1466** can be attached to the interior and/or exterior of the cuff frame **1468** using suitable fastening methods. For example, as shown in FIG. **14**B, the sleeve **1466** is positioned within the interior of the cuff frame **1468**, and the ends of the sleeve **1466** extend over and are fixed to proximal and distal ends of the cuff frame **1468** using suitable fastening methods (e.g., stitching, gluing, welding, etc.). In some embodiments, the proximal and distal ends of the cuff frame **1468** can be flared, and the sleeve **1466** can wrap around the flared ends to the exterior of the cuff frame **1468** such that the attachment can be sealed by the arterial walls when the aortic cuff **1464** is expanded to the expanded configuration in situ. The sleeve **1466** can have generally similar characteristics as the cover **106** described above. For example, the sleeve **1466** can be made from one or more substantially impermeable materials, such as Dacron® and PTFE, and can include ribs that can interface with arterial walls and/or endograft devices **102** (FIG. **14**B). The cuff frame **1468** can have generally similar characteristics as the integrated frame **104** described above. In other embodiments, the cuff frame **1468** can be made from individual zigzagged wire hoops like a Z-stent.

[0118] The sleeve **1466** and the cuff frame **1468** can have a substantially cylindrical shape. In some embodiments, the aortic cuff **1464** can include two channels to support superior portions **108** of endograft devices **102** (FIG. **14**B). For example, the channels can be formed by stitching the fabric of the sleeve **1466** together to divide the interior of the aortic cuff **1464**. Additionally, the sleeve **1466** and/or the cuff frame **1468** can have flared proximal and distal ends to form a stronger seal with adjacent arterial walls.

[0119] Referring to FIG. **14**B, the endograft devices **102** are deployed within the aortic cuff **1464** after the cuff **1464** has been at least substantially sealed against the aortic neck **60**. The superior portions **108** can mate with and substantially seal to the interior of the aortic cuff **1464**. The ribs **530** of the cover **106** can interface with the interior surface of the sleeve **1466** to further strengthen the seal. Additionally, the integrated frame **104** can further improve the seal between the endograft devices **102** and the aortic cuff **1464**. For example, the cross-section of the frame **104** in the expanded configuration can be slightly larger than an interior cross-section of the aortic cuff **1464**. As the endograft devices **102** are deployed within the aortic cuff **1464**, the radial forces from the expansion of the endograft devices **102** can strengthen the seal therebetween. Additionally, in some embodiments, the transition portion **324** of the endograft devices can mate with a complementary taper within the aortic cuff **1464**.

[0120] In some embodiments in accordance with the technology, the aortic cuff **1464** can include alignment aids, such as the alignment aids **734** described above with reference to FIGS. **7**A and **7**B, to facilitate positioning the endograft devices **102** within the aortic cuff **1464**. For example, the aortic cuff **1464** and the outer walls **112** of the endograft devices **102** can include orthogonal alignment aids that intersect to indicate the endograft devices **102** are properly aligned within the aortic cuff **1464**.

[0121] In additional embodiments, the aortic cuff **1464** can include anchors, such as the anchors **836** described above with reference to FIGS. **8**A and **8**B, to secure the to secure the system **1400** in situ. For example, the cuff frame **1468** can include anchors that project radially outwardly and engage adjacent arterial walls.

[0122] FIGS. **15**A and **15**B are schematic views of a three-part modular endograft system **1500** ("system **1500**") being deployed across the aneurysm **50** in accordance with an embodiment of the technology. The system **1500** can include the endograft devices **102** described with respect to the system **100** and the aortic cuff **1464** described above with reference to FIGS. **14**A and **14**B.

[0123] Referring to FIG. **15**A, the delivery system **40** can be inserted using a generally similar method as described above with reference to FIG. **9**A. In the embodiment shown in FIG. **15**A, however, the first catheter **42***a* and the first guidewire **44***a* can be inserted first to deliver the aortic cuff **1464** (FIG. **15**B) to the target site T. The aortic cuff **1464** can be deployed using a generally similar method as deploying the endograft devices **102** described above with reference to FIGS. **9**A and **9**B. The first guidewire **44***a* can be used to adjust the aortic cuff **1464** to a desired position in the aortic neck **60**.

[0124] As shown in FIG. **15**B, the endograft devices **102** can be deployed within the aortic cuff **1464**. The endograft devices **102** can be deployed using a substantially similar method as described with reference to FIG. **9**B. For example, the endograft devices **102** can be delivered through the first and second catheters **42** and positioned independently within the aortic cuff **1464** using the guidewires **44**. Similar to the method of deploying the superior portions **108** directly against the arterial walls described with reference to FIGS. **9**B and **13**B, here the outer walls of the superior portions **108** can at least partially interface with the interior surface of the aortic cuff **1464** such that the septal walls are aligned with each other to form the septum **120** (not visible). In some embodiments in accordance with the technology, the aortic cuff **1464** can include sections shaped to receive the endograft devices **102** and thereby ease alignment. In further embodiments, the first endograft device **102***a* can be anchored or otherwise secured to the aortic cuff **1464** before deployment such that only the second endograft device **102***b* must be positioned within the aortic cuff **1464**.

[0125] FIG. **16** is a schematic view of a modular endograft system **1600** ("system **1600**") being deployed across the aneurysm **50** in accordance with another embodiment of the technology. The system **1600** can be deployed using generally similar methods as the system **1500** described above with reference to FIGS. **15**A and **15**B. As shown in FIG. **16**, however, the superior portions **108** project above the aortic cuff **1464** such that the first end portions **118***a* provide additional structure for securing the endograft devices to the arterial walls of the aorta **52**. Additionally, the inferior portions **110** of the endograft devices **102** terminate within the aneurysm **50**. Therefore, the system **1600** further includes limbs (not shown), such as the limbs **1362** described above with reference to FIGS. **13**A-C, that connect to the inferior portions **110** and extend into the iliac arteries **56**. The catheters **42** can be used to adjust the length of the limbs to accommodate differing anatomies of the iliac arteries **56** and to maximize the fixation and sealing areas between the limbs and the arterial walls. Additionally, in some embodiments, the limbs can intersect (e.g., the limbs **1362** shown in FIG. **13**C) to strengthen the seal at the aortic neck **60** and decrease the likelihood of endoleaks. Similar to the four-part system **1300** described above, the five-part system **1600** can accommodate anatomical variations without requiring customized components.

[0126] In the embodiments illustrated in FIGS. **9**A, **9**B, **11**-**13**C, **15**A, **15**B and **16**, the aneurysm **50** is shown in the infrarenal portion of the aorta **52** because this is the most common site of an AAA. In other embodiments in accordance with the technology, the modular endograft systems **100**, **1000**, **1300**, **1500** and **1600** can be deployed across aneurysms **50** at different portions of the aorta **52** or in other vessels altogether. For example, in some embodiments, the aneurysm **50** can extend from the infrarenal portion of the aorta **52** into one or both of the common iliac arteries **56**. The inferior portions **110** or the limbs **1362** of the systems **100**, **1000**, **1300**, **1500** and **1600** can extend past the diseased, aneurysmal portion of the iliac arteries **56** without blocking blood flow to the internal iliac arteries **58**. In other embodiments, the systems **100**, **1000**, **1300**, **1500** and **1600** can be deployed across aneurysms **50** located in the supra renal portion of the aorta **52** with the fenestrations **1038** and/or the first end portions **118***a* positioned at the entrance of the renal arteries **54**. In further embodiments, the systems described above can be deployed across aneurysms in other portions of the vasculature that benefit from the use of a bifurcated, bi-luminal modular endograft system that can be independently positioned.

[0127] 4. Methods of Manufacturing

[0128] 4.1 Integrated Frame

[0129] Referring back to FIGS. **4**A and **4**B, the integrated frame **104** can be made by weaving or braiding one continuous wire **426** in a pattern along a cylindrical mandrel. In some embodiments, the wire **426** is woven with a one over and one under pattern. In other embodiments, the wire **426** is woven with a two over and one under pattern, another integrated pattern, and/or a pattern that varies over the length of the frame **104**. The intersections of the wire **426** can remain

unbound to increase flexibility of the frame **104**. The wire **426** can form the loops **428** to change direction and continue the pattern of intersecting wires **426**. As described above, the number of loops **428** at each end portion **118** and the braid angle θ can be selected based on the diameter of the wire **426** and the desired properties of the frame **104**.

[0130] The wire **426** can be removed from the mandrel after it is braided into the frame **104** and formed into a desired shape (e.g., the endograft devices **102** shown above). The frame **104** can then be heated to a shape-setting temperature specified for the wire material (e.g., Nitinol), and subsequently quenched. Optionally, the frame **104** can be annealed to increase the strength of the frame **104**. The mandrel can be cylindrical or have the shape of the frame **104** such that the wire **426** remains on the mandrel during heat treatment. In further embodiments, the frame **104** can be manufactured using other suitable methods for shaping resilient biocompatible materials.

[0131] 4.2 Covers and Coatings

[0132] Referring to FIGS. **5**A-C, the cover **106** can be made by shaping a substantially non-permeable cover material, such as Dacron®, PTFE, and/or other suitable biocompatible materials. The cover **106** can be formed by first placing the cover material over a mandrel. The mandrel can include thin grooves that can correspond to the desired geometry of the ribs **530** on the cover **106**. A wire or thread can be wrapped over the cover material and into the grooves to corrugate the cover material. The cover material can then be heated on the mandrel until the ribs **530** are formed and the cover **106** is substantially non-permeable. In some embodiments, the superior and inferior termini **531***a* and **531***b* of the cover **106** can be shaped to facilitate attaching the cover **106** to a frame (e.g. the frame **104** shown in FIGS. **4**A and **4**B) and prevent the cover **106** from wrinkling at end portions (e.g., the end portions **118** shown in FIGS. **1**A and **1**B) during constriction. For example, the superior and inferior termini **531***a* and **531***b* can be zigzagged as shown in FIGS. **5**A and **5**B, scalloped, or otherwise shaped to limit wrinkling of the cover on the frame.

[0133] In other embodiments in accordance with the technology, coating layers can be used in place of or in conjunction with the cover **106**. FIGS. **17**A-E are views of coating layers being applied to an integrated frame **1704** ("frame **1704**") in accordance with embodiments of the technology. The frame **1704** has generally similar features as the frame **104** described above. For example, the frame **1704** can be made from the braided wire **426**.

[0134] Referring to FIG. **17**A, the frame **1704** is positioned over a mandrel **80** in the expanded configuration. As shown in FIG. **17**A, a first coating layer **1770** can be wrapped onto the frame **1704**. The first coating layer **1770** can be a single or double layer of unsintered tape that can be approximately 0.0005" thick and made from PTFE. In other embodiments, the first coating layer **1770** can have a different thickness and/or the first coating layer **1770** can be made from another suitable coating material.

[0135] Once the first coating layer **1770** is applied over the frame **1704**, the first coating layer **1770** and the frame **1704** can be heated on the mandrel **80** in an oven. For example, the first coating layer **1770** and the frame **1704** can be heated for less than thirty minutes in a 370° C. oven. After heating, the coated frame **1704** is removed from the mandrel **80** and extended and contracted from the low-profile configuration to the expanded configuration to ensure the first coating layer **1770** properly adhered to the frame **1704** during heat treatment.

[0136] As shown in FIG. **17**B, a second coating material **1772** is placed over a narrower, second mandrel **82**. The second coating material **1772** can be extended a distance equivalent to the length of the frame **1704** in the low-profile configuration. Referring to FIG. **17**C, the second coating material **1772** is contracted to the length of the frame **1704** in the expanded configuration. This contraction can form small ribs **1730** in the second coating material **1772**. The ribs **1730** can be generally similar to the ribs **530** described above with reference to FIGS. **5**A-C, but they are on the interior of the frame **1704**. The ribs **1730** prevent the second coating material **1772** from wrinkling or bunching when the subsequently attached frame **1704** flexes or bends and thereby reduce the likelihood of thrombotic problems within the lumen.

[0137] As shown in FIG. **17**D, the coated frame **1704** is then extended to the low-profile configuration and placed over the extended second coating material **1772** on the second mandrel **82**. Each diamond opening along the frame **1704** can be spot welded using a welding device **84**. Then, the frame **1704** is removed from the second mandrel **82** and extended and contracted from the low-profile configuration to the expanded configuration to ensure that the first and second coating layers **1770** and **1772** have adequately adhered to the frame **1704**. Additionally, the proximal and distal ends of the frame **1704** are verified to ensure that the first and second coating layers **1770** and **1772** have properly adhered to the frame **1704**. If necessary, tacking can be performed and the edges can be trimmed to form a dual coated endograft device **1702** shown in FIG. **17**E.

[0138] From the foregoing, it will be appreciated that specific embodiments of the invention have been described herein for purposes of illustration, but that various modifications may be made without deviating from the spirit and scope of the technology. For example, the embodiments illustrated in FIGS. **1**A-**16** include covers **106** that extend over the exterior of the integrated frames **104**. However, other embodiments of the technology can include covers **106** that are attached to the interior of the integrated frame **104** and/or are formed integrally with the frame **104**. Certain aspects of the new technology described in the context of particular embodiments may be combined or eliminated in other embodiments. For example, in the embodiments illustrated above, each endograft device (e.g., **102**, **1002**) includes a singular lumen **116**. However, the endograft devices can include additional lumens that transverse, bisect, and/or otherwise communicate with the lumen **116** to accommodate the vasculature. For example, the endograft devices can include lumens that extend into the renal arteries, the internal iliac arteries, and/or other arteries. Further, while advantages associated with certain embodiments of the technology have been described in the context of those embodiments, other embodiments may also exhibit such advantages, and not all embodiments need necessarily exhibit such advantages to fall within the scope of the technology. Accordingly, the disclosure and associated technology can encompass other embodiments not expressly shown or described herein.

I/We claim:

1. A modular endograft system, comprising:

a cuff having a proximal end portion and a distal end portion;

a first endograft device having a first superior portion, a first inferior portion, and a first lumen through the first superior and inferior portions, wherein the first superior portion has a first outer wall and a first septal wall;

a second endograft device having a second superior portion, a second inferior portion, and a second lumen through the second superior and inferior portions, wherein the second superior portion has a second outer wall and a second septal wall; and

wherein the first and second endograft devices are configured such that the first and second outer walls of the first and second endograft devices press against the cuff and the first and second septal walls exert opposing forces toward one another to fix the cuff to the first and second lumens.

2. The modular endograft system of claim **1** wherein the cuff has a substantially circular cross section and at least one of the proximal and distal end portions of the cuff is flared radially outward.

3. The modular endograft system of claim **1** wherein the cuff includes a cuff frame having a proximal terminus and a distal terminus and a sleeve radially inward from the cuff frame, wherein the sleeve extends over the proximal and distal termini of the cuff frame.

4. The modular endograft system of claim **1** wherein:

the cuff includes a sleeve and a cuff frame at least partially enclosing the sleeve;

the sleeve bifurcates the cuff into a first cuff lumen and a second cuff lumen; and

the first superior portion of the first endograft device contacts the first cuff lumen and the second superior portion of the second endograft device contacts the second cuff lumen.

5. The modular endograft system of claim **1** wherein the first and second superior portions of the first and second endograft devices, respectively, project distally beyond the distal end portion of the cuff such that portions of the first and second outer walls of the endograft devices affix to an adjacent vessel wall.

6. The modular endograft system of claim **1** wherein:

the first superior portion of the first endograft device includes a convexly curved first outer wall having a first radius and a convexly curved first septal wall having a second radius greater than the first radius such that the first superior portion has a D-like cross-section;

the second superior portion of the second endograft device includes a convexly curved second outer wall having the first radius and a convexly curved second septal wall having the second radius such that the second superior portion has a D-like cross-section;

wherein the first and second outer walls press against the cuff device in an expanded configuration; and

the first and second septal walls press against each other in the expanded configuration and form a septum, and the convex curvature of the first and second septal walls results in a substantially uniform distribution of pressure along the septum.

7. The modular endograft system of claim **6** wherein:

the first septal wall and the first outer wall are joined at curved corners;

the second septal wall and the second outer wall are joined at curved corners; and

the curved corners have a radius of curvature less than the first radius, and wherein the curved corners form an angle from approximately 60° to approximately 100° at the septum in the expanded configuration.

8. The modular endograft system of claim **1**, further comprising at least one anchor coupled to an exterior of the cuff, wherein the anchor protrudes radially from the cuff in an expanded configuration and constricts in a low-profile configuration.

9. The modular endograft system of claim **1** wherein the first and second endograft devices include at least one anchor at the first and second superior portions, wherein the anchor protrudes radially from the superior portions in an expanded configuration.

10. The modular endograft system of claim **1** wherein:

the first endograft device includes a first alignment aid at the first septal wall; and

the second endograft device includes a second alignment aid at the second septal wall, wherein the second alignment aid crosses the first alignment aid when the first and second septal walls of the first and second endograft devices, respectively, oppose one another such the first and second septal walls form a septum.

11. The modular endograft system of claim **1** wherein:

the cuff includes a first cuff alignment aid and a second cuff alignment aid;

the first endograft device includes a first alignment aid at the first outer wall, wherein the first alignment aid crosses the first cuff alignment aid when the first outer wall opposes the cuff; and

the second endograft device includes a second alignment aid at the second outer wall, wherein the second alignment aid crosses the second cuff alignment aid when the second outer wall opposes the cuff.

12. The modular endograft system of claim **1** wherein:

the first endograft device further includes a first transition portion between the first superior portion and the first inferior portion, the first transition portion tapering the first lumen from a first cross-sectional dimension at the superior portion to a second cross-sectional dimension less than the first cross-sectional dimension at the inferior portion, wherein the transitional portion is configured to maintain substantially laminar blood flow through the first lumen in an expanded configuration;

the second endograft device further includes a second transition portion between the second superior portion and the second inferior portion, the second transition portion tapering the second lumen from the first cross-sectional dimension at the second superior portion to the second cross-sectional dimension at the second inferior portion, wherein the second transitional portion is configured to maintain substantially laminar blood flow through the second lumen in the expanded configuration; and

the cuff further includes a transitional cuff portion at the proximate end portion, the tapered cuff portion being substantially conformal to the first and second tapered transitional portions.

13. The modular endograft system of claim **1** wherein the first and second endograft devices each include a frame having a superior terminus, an inferior terminus, and a continuous wire woven in a braid, the wire crossing itself at a braid angle, and the wire reversing direction at the superior terminus of the frame to form a first plurality of loops and reversing direction at the inferior terminus of the frame to form a second plurality of loops.

14. The modular endograft system of claim **13** wherein the first and second endograft devices each further include a cover over at least a portion of the frame, the cover having circumferential ribs protruding radially from the frame such that opposing circumferential ribs of the first and second septal walls mate in the expanded configuration.

15. A modular endograft system, comprising:

a cuff having a cuff frame and a sleeve attached to the cuff frame, wherein the cuff frame and the sleeve have a proximal end portion and a distal end portion;

a first endograft device having a first frame, a first cover attached to the first frame, and a first lumen within the first cover, wherein the first frame and the first cover have a first superior portion and a first inferior portion, and the first superior portion has a convexly curved first outer wall and a first septal wall;

a second endograft device having an integrated second frame ("second frame"), a second cover attached to the second frame, and a second lumen within the second cover, wherein the second frame and the second cover have a second superior portion and a second inferior portion, the second superior portion having a convexly curved second outer wall and a second septal wall; and

wherein the first and second endograft devices are configured to be extended into a low-profile configuration with a first cross-sectional dimension and expand to an expanded profile configuration with a second cross-sectional dimension greater than the first cross-sectional dimension such that in the expanded configuration the first and second septal walls are urged toward each other and form a septum between the first and second lumens and the first and second outer walls press against an interior of the cuff.

16. The modular endograft system of claim **15** wherein at least one of the proximal and distal end portions of the cuff frame is flared radially outward.

17. The modular endograft system of claim **16** wherein the sleeve extends over proximal and distal ends of the cuff frame such that the sleeve attaches to the cuff frame when the cuff presses against vessel walls.

18. The modular endograft system of claim **15** wherein the sleeve includes a first cuff lumen configured to receive the first endograft device and a second cuff lumen configured to receive the second endograft device.

19. The modular endograft system of claim **15** wherein the first and second endograft devices are staggered longitudinally relative to the first and second lumens such that the first superior portion includes a free end portion projecting distally beyond the second superior portion of the second endograft device.

20. The modular endograft system of claim **15** wherein the first and second frames each include a superior terminus, an inferior terminus, and a continuous wire woven in a braid, the wire crossing itself at a braid angle, and the wire reversing direction at the superior terminus to form a first plurality of loops and reversing direction at the inferior terminus to form a second plurality of loops

21. The modular endograft system of claim **20** wherein the braid angle is from approximately 30° to approximately 45°.

22. The modular endograft system of claim **20** wherein the wire has a diameter from approximately 0.0070 inch to approximately 0.0140 inch and the first plurality of loops includes no more than eight loops and the second plurality of loops includes no more than eight loops.

23. The modular endograft system of claim **15** wherein the first and second covers extend over the first and second frames, and wherein the first and second covers limit at least one of radial expansion and longitudinal contraction of the first and second frames, respectively, in the expanded configuration.

24. The modular endograft system of claim **15** wherein:

at least a portion of the sleeve is radially inward from the cuff frame, and wherein the sleeve includes ribs protruding radially inward from the cuff frame;

the first cover includes first ribs protruding radially from the first frame in the dilated configuration, the first ribs being extendable longitudinally in the low-profile configuration;

the second cover includes second ribs protruding radially from the second frame in the dilated configuration, the second ribs being extendable longitudinally in the low-profile configuration; and

wherein the first and second ribs interface with the ribs of the sleeve in the expanded configuration.

25. The modular endograft system of claim **15**, further comprising at least one anchor protruding radially outward from an exterior of the cuff.

26. The modular endograft system of claim **15** wherein the first and second superior portions include alignment aids on opposing first and second septal walls, the alignment aids being configured to cross one another when the first and second septal walls form the septum.

27. A method of repairing an aneurysm in a primary blood vessel before a bifurcation into a first blood vessel and a second blood vessel, comprising:

advancing a cuff through the first blood vessel to a target site in the primary blood vessel before the aneurysm, the cuff having a substantially circular cross-section;

deploying the cuff at the target site such that the cuff presses against a vessel wall of the primary blood vessel;

advancing a first endograft device through the first blood vessel to a target site in the common blood vessel before the aneurysm;

advancing a second endograft device through the second blood vessel to the target site;

deploying the first and second endograft devices at the target site such that the first and second endograft devices expand to an expanded configuration via inherent spring forces in the first and second endograft devices such that first and second septal walls of the first and second endograft devices, respectively, press toward each other and form a septum between the first and second endograft devices, and wherein first and second outer walls of the first and second endograft devices, respectively, sealably press against an interior surface of the cuff; and

wherein the first and second endograft devices are positioned independently of one another.

28. The method of claim **27** wherein positioning the first and second endograft devices independently comprises:

positioning a first superior portion of the first endograft device in a first desired position within the cuff; and

positioning a second superior portion of the second endograft device in a second desired position within the cuff, wherein the first desired position is longitudinally offset from the second desired position along the cuff.

29. The method of claim **27**, further comprising:
loading the first endograft device in a first catheter, wherein the first endograft device is extended in a low-profile configuration;
loading the second endograft device in a second catheter, wherein the second endograft device is extended in a low-profile configuration;
percutaneously introducing the first endograft device into the first blood vessel; and
percutaneously introducing the second endograft device into the second blood vessel.

30. The method of claim **29** wherein the first and second endograft devices have a cross-sectional dimension no less than 20 mm in the dilated configuration, and wherein the first and second catheters are no larger than 12 F.

31. The method of claim **27** wherein the first endograft device includes a first alignment aid at the first septal wall and the second endograft device includes a second alignment aid at the second septal wall, the first and second alignment aids comprising a radiopaque material, and wherein deploying the first and second endograft devices further includes:
radiographically positioning the first and second alignment aids such that the first and second alignment aids oppose one another;
viewing the first and second alignment aids in the orthogonal plane; and
crossing the first and second alignment aids such that the first and second septal walls can form the septum.

32. The method of claim **31** wherein the first and second alignment aids diagonally cross the first and second septal walls, and wherein the first and second alignment aids form an "X" indicator at the septum.

33. The method of claim **27** wherein the cuff includes a cuff frame and an internal sleeve having ribs protruding radially inward from the cuff frame, the first and second endograft devices each include a cover having a plurality of circumferential ribs at least at the first and second septal and outer walls of the first and second endograft devices, respectively, and wherein deploying the first and second endograft devices further comprises:
interfacing the circumferential ribs at the first septal wall with the circumferential ribs at the second septal wall; and
interfacing the circumferential ribs at the first and second outer walls with the ribs of the internal sleeve.

34. The method of claim **27** wherein the first and second endograft devices each include a cover attached over at least a portion of a frame, and deploying the first and second endograft devices further comprises restricting radial expansion of the frames with the covers.

35. The method of claim **27** wherein the primary blood vessel is an aorta and the first blood vessel is a first common iliac artery and the second blood vessel is a second common iliac artery, and wherein deploying the cuff and the first and second endograft devices further includes:
at least substantially sealing the cuff with an arterial wall of the aorta;
at least substantially sealing the first inferior portion with an interior arterial wall of the first common iliac artery; and
at least substantially sealing the second inferior portion with an interior arterial wall of the second common iliac artery.

36. The method of claim **27** wherein the primary blood vessel includes a third blood vessel and a fourth blood vessel branching from the primary blood vessel before the aneurysm, the third and fourth blood vessels being longitudinally offset from one another relative to the primary blood vessel, and wherein:
the first endograft device has a first braided frame and a first cover coupled to the first braided frame, the first cover having a first inferior terminus and a first superior terminus, and the first braided frame having a first end extending distally beyond the first superior terminus of the first cover and second end extending proximally beyond the first inferior terminus of the first cover, wherein the first and second end portions include openings through which blood can flow laterally relative to a longitudinal axis of the first lumen;
the second endograft device has a second braided frame and a second cover attached to the second braided frame, the second cover having a second inferior terminus and a second superior terminus, and the second braided frame having a first end extending distally beyond the second superior terminus of the second cover and a second end extending proximally beyond the second inferior terminus of the second cover, wherein the first and second end portions include openings through which blood can flow laterally relative to a longitudinal axis of the second lumen;
deploying the cuff comprises positioning the cuff proximal to the third and fourth blood vessels; and
deploying the first and second endograft devices comprises positioning the first end portion of the first frame at the entrance of the third blood vessel and positioning the first end portion of the second frame at the entrance of the fourth blood vessel such that the third and fourth blood vessels are in fluid communication with blood flow through the first and the second lumens.

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