<table>
<thead>
<tr>
<th>CPT® Code</th>
<th>Description</th>
<th>2020 Work RVUs</th>
<th>Applicable Modifiers*</th>
<th>Case Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Surgical Arterial Access</td>
<td></td>
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<tr>
<td>+34713</td>
<td>Perc access and closure of femoral artery</td>
<td>2.50</td>
<td>-50 -51 -62</td>
<td>-80 -82 -AS</td>
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<tr>
<td>+34714</td>
<td>Open femoral exposure with creation of conduit</td>
<td>5.25</td>
<td>-50 -62 -80</td>
<td>-82 -AS</td>
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<tr>
<td>+34812</td>
<td>Open femoral exposure</td>
<td>4.13</td>
<td>-50 -62 -80</td>
<td>-82 -AS</td>
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<tr>
<td>+34820</td>
<td>Open iliac exposure</td>
<td>7.00</td>
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<td>-82 -AS</td>
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<tr>
<td>+34833</td>
<td>Open iliac exposure with creation of conduit</td>
<td>8.16</td>
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<td>-82 -AS</td>
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<tr>
<td>+34834</td>
<td>Open brachial exposure</td>
<td>2.65</td>
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<td>-82 -AS</td>
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<tr>
<td>Placement of Wires, Catheters, Sheaths</td>
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<tr>
<td>36200</td>
<td>Catheter/sheath placement into aorta; nonselective</td>
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<td>36215</td>
<td>Catheter/sheath placement; selective, first order</td>
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<td>36216</td>
<td>Catheter/sheath placement; selective, second order</td>
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<td>36217</td>
<td>Catheter/sheath placement; selective, third order</td>
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<tr>
<td>Placement and Deployment of Endoluminal Stent Graft</td>
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<td>33880</td>
<td>Endo TAA repair with coverage of LSA, to celiac if req.</td>
<td>34.58</td>
<td>-50 -51 -62 -80</td>
<td>-82 -AS</td>
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<tr>
<td>75956-26</td>
<td>Rad. S&amp;I: endovascular TAA repair</td>
<td>7.00</td>
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<tr>
<td>33881</td>
<td>Endo TAA repair without coverage of LSA, to celiac if req.</td>
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<td>-82 -AS</td>
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<tr>
<td>75957-26</td>
<td>Rad. S&amp;I: endovascular TAA repair</td>
<td>6.00</td>
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<tr>
<td>75958-26</td>
<td>Rad. S&amp;I: extension prosthesis</td>
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<td>+33884</td>
<td>Ext. Proximal each additional</td>
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<td>75958-26</td>
<td>Rad. S&amp;I: extension prosthesis</td>
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<td>33886</td>
<td>Ext. Distal, delayed after initial endovascular repair</td>
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<td>75959-26</td>
<td>Rad. S&amp;I: extension prosthesis, delayed placement</td>
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<td>Ancillary Procedures</td>
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<td>+37252</td>
<td>IVUS noncoronary, initial vessel</td>
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<td>IVUS noncoronary, additional vessel</td>
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<td>37242</td>
<td>Art. embolization or coiling (non-hemorrhage or tumor)</td>
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<td>62272</td>
<td>Spinal puncture, therapeutic (Lumbar drain)</td>
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<td>37236</td>
<td>Non-coronary arterial stent, initial artery</td>
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<tr>
<td>33889</td>
<td>Transposition; open sub-c to carotid, neck incision with TAA</td>
<td>15.92</td>
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<td>33891</td>
<td>Carotid-carotid bypass graft, in conjunction with TAA</td>
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<td>34712</td>
<td>Trans cath delivery of enhanced fixation device</td>
<td>12.00</td>
<td>-51 -62 -80</td>
<td>-82 -AS</td>
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</tbody>
</table>

* Other coding modifiers may apply. + Add-on code; list in addition to primary procedure. These suggestions do not replace seeking coding advice from the payer and/or your coding staff, the provider of services is ultimately responsible for correct coding.
The Valiant Navion™ thoracic stent graft system is contraindicated in the following patient populations:

- Patients who have a condition that threatens to infect the graft
- Patients who are sensitive to or have allergies to the device materials

Adverse events or complications associated with the use of the Valiant Navion thoracic stent graft system that may occur or require intervention include, but are not limited to:

- Access failure
- Access site complications (for example: spasm, trauma, bleeding, rupture, dissection)
- Adynamic ileus
- Allergic reaction (to contrast, antiplatelet therapy, stent graft material)
- Anaphylaxis
- Aneurysm rupture
- Angina
- Aortic expansion (for example: aneurysm, false lumen)
- Aortic valve damage
- Aortic vessel rupture
- Arrhythmia
- Arterial stenosis
- Atelectasis
- Balloon rupture
- Blindness
- Bowel ischemia
- Bowel necrosis
- Branch vessel occlusion
- Catheter breakage
- Cerebrovascular accident
- Change in mental status
- Coagulopathy
- Congestive heart failure
- Conversion to surgical repair
- Damage to the vessel
- Death
- Deployment dysfunction
- Deployment failure
- Deployment misplacement
- Dissection
- Embolism
- Endoleak
- Excessive or inappropriate radiation exposure
- Extrusion/erosion
- Failure to deliver the stent graft
- Femoral neuropathy
- Fistula
- FreeFlo and CoveredSeal configuration
- Genitourinary complications
- Hematoma
- Hemorrhage/bleeding
- Hypertension/hypertension
- Infection or fever
- Insertion or removal difficulty
- Intercostal pain
- Intramural hematoma
- Leg edema/foot edema
- Loss of potency
- Lymphohoele
- Myocardial infarction
- Neck enlargement
- Nerve injury
- Neuroptopathy
- Occlusion – Venous or Arterial
- Pain
- Patient reaction at catheter insertion site
- Paralysis
- Paraparesis
- Paraplegia
- Paresthesia
- Perforation of the false lumen
- Peripheral ischemia
- Peripheral nerve injury
- Pneumonia
- Postimplant syndrome
- Post-procedural bleeding
- Procedural bleeding
- Prosthesis dilatation
- Prosthesis infection
- Prosthesis rupture
- Pseudoaneurysm
- Pulmonary edema
- Reaction to anesthesia
- Renal failure
- Renal insufficiency
- Reoperation
- Respiratory depression or failure
- Retrograde type A dissection
- Sepsis
- Seroma
- Sexual dysfunction
- Shock
- Spinal neurological deficit
- Stenosis
- Stent graft migration
- Stent graft misplacement
- Stent graft rupture
- Stent graft twisting or kinking
- Transient ischemic attack
- Thrombosis
- Tissue necrosis
- Vascular ischemia
- Vascular trauma
- Wound dehiscence
- Wound healing complications
- Wound infection

Never use a balloon when treating a dissection.

Please refer to the product Instructions for Use for details.

MRI Safety and Compatibility

MRI may be used on the Valiant Navion thoracic stent graft only under specific conditions. It can be scanned safely in both 1.5T & 3.0T MR systems under certain conditions as described in the product Instructions for Use. For additional information regarding MRI please refer to the product Instructions for Use.

Adverse Events

Adverse events or complications associated with the use of the Valiant Navion thoracic stent graft system that may occur or require intervention include, but are not limited to:

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Please refer to the product Instructions for Use for more information regarding indications, warnings, precautions, contraindications and adverse events.

CAUTION: Federal (USA) law restricts this device to sale by or on the order of a physician.