Medical Policy

Wireless Pressure Sensors in Endovascular Aneurysm Repair

Table of Contents

- Policy: Commercial
- Policy: Medicare
- Authorization Information
- Coding Information
- Description
- Policy History
- Information Pertaining to All Policies
- References

Policy Number: 306
BCBSA Reference Number: 7.01.111

Related Policies

- Cardiac Hemodynamic Monitoring for the Management of Heart Failure in the Outpatient Setting #287
- Endovascular Grafts for Abdominal Aortic Aneurysms #098
- Endovascular Stent Grafts for Thoracic Aortic Aneurysms or Dissections #233

Policy

Commercial Members: Managed Care (HMO and POS), PPO, and Indemnity Medicare HMO BlueSM and Medicare PPO BlueSM Members

Wireless pressure sensors in the management (intraoperative and/or postoperative) of patients having endovascular aneurysm repair are INVESTIGATIONAL.

Prior Authorization Information

Pre-service approval is required for all inpatient services for all products. See below for situations where prior authorization may be required or may not be required for outpatient services.

Yes indicates that prior authorization is required.
No indicates that prior authorization is not required.
N/A indicates that this service is primarily performed in an inpatient setting.

<table>
<thead>
<tr>
<th>Outpatient</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Commercial Managed Care (HMO and POS)</td>
<td>This is not a covered service.</td>
</tr>
<tr>
<td>Commercial PPO and Indemnity</td>
<td>This is not a covered service.</td>
</tr>
<tr>
<td>Medicare HMO BlueSM</td>
<td>This is not a covered service.</td>
</tr>
<tr>
<td>Medicare PPO BlueSM</td>
<td>This is not a covered service.</td>
</tr>
</tbody>
</table>

CPT Codes / HCPCS Codes / ICD-9 Codes

The following codes are included below for informational purposes. Inclusion or exclusion of a code does not constitute or imply member coverage or provider reimbursement. Please refer to the member’s
contract benefits in effect at the time of service to determine coverage or non-coverage as it applies to an individual member.

Providers should report all services using the most up-to-date industry-standard procedure, revenue, and diagnosis codes, including modifiers where applicable.

CPT Codes

<table>
<thead>
<tr>
<th>CPT codes</th>
<th>Code Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>34806</td>
<td>Transcatheter placement of wireless physiologic sensor in aneurysmal sac during endovascular repair, including radiological supervision and interpretation, instrument calibration, and collection of pressure data (List separately in addition to code for primary procedure)</td>
</tr>
<tr>
<td>93982</td>
<td>Noninvasive physiologic study of implanted wireless pressure sensor in aneurysmal sac following endovascular repair, complete study including recording, analysis of pressure and waveform tracings, interpretation and report</td>
</tr>
</tbody>
</table>

ICD-9 Diagnosis Codes

Investigational for all diagnoses.

Description

The goal of abdominal aortic aneurysm (AAA) repair is to reduce pressure in the aneurysm sac and thus prevent rupture. Failure to exclude the aneurysm completely from the systemic circulation results in continued pressurization. An endoleak (persistent perfusion of the aneurysmal sac) may be primary (within the first 30 days post-op) or secondary (after 30 days post-op). Endoleaks are reported to occur in 10–50% of cases, and there are 5 types of endoleaks (I-V).

Wireless sensors implanted in an aortic aneurysm sac after endovascular repair are being investigated to measure post-procedural pressure. These implanted devices use various mechanisms to wirelessly transmit pressure readings to devices for measuring and recording pressure and have the potential to improve outcomes for patients who have had endovascular repair. It is thought that low pressures may correlate with positive prognoses and high pressures may indicate the need for revision.

It is also argued that wireless pressure sensors may change the need for or the frequency of monitoring of the aneurysm sac using contrast-enhanced computed tomography (CT) scans. They may improve postoperative monitoring.

An example of a wireless pressure sensor in endovascular aneurysm repair is the CardioMEMS EndoSure™ from CardioMEMS, Inc. All wireless pressure sensors in endovascular aneurysm repair are considered investigational regardless of the commercial name, the manufacturer or FDA approval status.

Summary

Data are currently insufficient to indicate if use of this device improves clinical outcomes. The accuracy of the device in those with different types of endoleaks needs to be determined with larger numbers of patients. Also, the performance over time needs to be addressed. Work is also needed to determine the type and number of devices that might best be used in monitoring given that sac compartmentalization might lead to a pressure-sensing device missing an endoleak. It also is not known whether there might be important long-term complications from this implanted device. Furthermore, the extent to which the device can reduce imaging requirements following endovascular aneurysm repair (via direct comparison with CT) is undetermined. The evidence to date, which consists of small case series, is insufficient to permit conclusions concerning the effect of this device on health outcomes. Therefore, the use of wireless pressure sensors in detecting endoleaks in aneurysm repair is considered investigational.
Policy History

<table>
<thead>
<tr>
<th>Date</th>
<th>Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>1/19/2011</td>
<td>New policy effective 1/19/2011 describing ongoing non-coverage.</td>
</tr>
</tbody>
</table>

Information Pertaining to All Blue Cross Blue Shield Medical Policies
Click on any of the following terms to access the relevant information:
- Medical Policy Terms of Use
- Managed Care Guidelines
- Indemnity/PPO Guidelines
- Clinical Exception Process
- Medical Technology Assessment Guidelines

References