



June 13, 2020

CHAD BOERST  
 ABMRC LLC  
 239 FAIRCHILD ST  
 CHARLESTON, SC 29492

**DCN Number:20112003000000**

Manufacturer Name	Product Name	Model Number	Assigned HCPCS Code(s)
ABMRC LLC	BIWAZE COUGH SYSTEM	PRTN-2037351818-2 0107	E0482
ABMRC LLC	SIZE 0 PATIENT CIRCUIT	PRTN-2037351818-2 0114	A7020
ABMRC LLC	SIZE 1 PATIENT CIRCUIT	DDOC-1789753845-2 1085	A7020
ABMRC LLC	SIZE 2 PATIENT CIRCUIT	DDOC-1789753845-2 1086	A7020
ABMRC LLC	SIZE 3 PATIENT CIRCUIT	DDOC-1789753845-2 1087	A7020
ABMRC LLC	SIZE 4 PATIENT CIRCUIT	DDOC-1789753845-2 1088	A7020
ABMRC LLC	SIZE 5 PATIENT CIRCUIT	DDOC-1789753845-2 1089	A7020
ABMRC LLC	TRACH ADAPTOR PATIENT CIRCUIT	DDOC-1789753845-2 1090	A7020

Dear CHAD BOERST,

The Pricing, Data Analysis, and Coding (PDAC) Contractor has reviewed the product(s) listed above and has approved the listed Healthcare Common Procedure Coding System (HCPCS)

code(s) for billing the four Durable Medical Equipment Medicare Administrative Contractors (DME MACs).

The PDAC Contractor provides coding assistance to manufacturers to ensure proper coding of Durable Medical Equipment, Prosthetics, Orthotics, and Supplies (DMEPOS). The PDAC publishes coding decisions based on the coding guidelines established by the Local Coverage Determinations (LCDs) and associated Policy Articles and any related Advisory Articles established by the DME MACs. All products submitted to the PDAC for a coding verification review are examined by coders and professionals following a formal, standardized process.

Based on this review and application of DME MAC policy, the HCPCS code(s) listed below should be used when billing the DME MACs:

E0482 COUGH STIMULATING DEVICE, ALTERNATING POSITIVE AND NEGATIVE AIRWAY PRESSURE

A7020 INTERFACE FOR COUGH STIMULATING DEVICE, INCLUDES ALL COMPONENTS, REPLACEMENT ONLY

If you disagree with this decision, you may request a reconsideration within 45 days of the letter's date and provide evidence to substantiate a reconsideration of PDAC's original coding determination. To request a reconsideration, complete the Reconsideration Request form located on the PDAC website at [www.dmepdac.com](http://www.dmepdac.com). If your request for a reconsideration is made after the 45-day time frame, it will require a new application and documentation to support the request.

It is the responsibility of manufacturers and distributors to notify the PDAC immediately of any changes involving their products, as listed on the Product Classification List (PCL) on the Durable Medical Equipment Coding System (DMECS). Further information for requesting updates to the PCL can be found on the PDAC website at [www.dmepdac.com](http://www.dmepdac.com). It is also the responsibility of manufacturers and distributors to assure their websites and product marketing materials accurately reflect the product reviewed by the PDAC and the coding decision assigned.

An assignment of the HCPCS code(s) to product(s) is not an approval or endorsement of the product(s) by Medicare or Palmetto GBA; nor does it imply or guarantee claim reimbursement or coverage.

If you have questions, please contact the PDAC HCPCS Helpline at (877) 735-1326 during the hours of 9:30 a.m. to 5:00 p.m. ET, Monday through Friday. You may also visit our [website](#) to chat with one of our representatives or select the Contact Us button at the top of the page for

email, FAX or postal mail information.

Sincerely,

Pricing, Data Analysis, and Coding Contract (PDAC)  
Palmetto GBA, LLC  
[www.dmepdac.com](http://www.dmepdac.com)