



July 11, 2025

LASSE MADSEN
LEVITATE TECHNOLOGY APS
MARIELUNDVEJ 28 ST
HERLEV, DENMARK, 2730

Document Control Number (DCN): 25106C25100000

Manufacturer Name	Product Name	Model Number	Assigned HCPCS Code(s)
LEVITATE TECHNOLOGY APS	LEVITATE BLADE	LTA00510-15	L5999
LEVITATE TECHNOLOGY APS	LEVITATE BLADE	LTA006101-18	L5999

Dear LASSE MADSEN,

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:

THIS ITEM HAS BEEN CORRECTED TO BE DESCRIBED AS L5999 - LOWER EXTREMITY PROSTHESIS, NOT OTHERWISE SPECIFIED.

UPON RE-REVIEW, THE LEVITATE BLADE DOES NOT INCLUDE A HEEL COMPONENT FEATURE WHICH IS REQUIRED TO BE DESCRIBED BY L5981 AS DESCRIBED ITALICS BELOW FROM THE LOWER LIMB PROSTHESES - POLICY ARTICLE A52496:

*L5981 describes a product that can be used for either endoskeletal or exoskeletal lower limb construction. The Flex Walk has an energy storing J-shaped keel design. **Heel component is attached to the J-shaped keel section.** The Flex Walk J-shaped keel design proximally terminates at a nonadjustable fixed height determined and modified by the prosthetic foot manufacturer. L5981 includes foot cover.*

THERE ARE NO EXISTING HCPCS CODES THAT DESCRIBE THE FEATURES AND FUNCTION OF THIS PRODUCT, THEREFORE, NOC CODE L5999 ADEUQUATLEY DESCRIBES YOUR PRODUCT.

If you disagree with this decision, you may request a reconsideration within 45 calendar days of the Coding Verification 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. If your request for a reconsideration of PDAC's coding determination is made after the 45 calendar 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. 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 (PDAC)
Palmetto GBA, LLC
www.dmepdac.com