



July 18, 2019

JEFFREY BAKER
MEDI USA LP
6481 FRANZ WARNER PARKWAY
WHITSETT, NC 27377

DCN Number:19182C28100000

Manufacturer Name	Product Name	Model Number	Assigned HCPCS Code(s)
MEDI USA LP	M.3 SOFT OA	KL01XXX	L1843
MEDI USA LP	M.3 SOFT OA	KL01XXX	L1851

Dear JEFFREY BAKER,

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:

- L1843 KNEE ORTHOSIS, SINGLE UPRIGHT, THIGH AND CALF, WITH ADJUSTABLE FLEXION AND EXTENSION JOINT (UNICENTRIC OR POLYCENTRIC), MEDIAL-LATERAL AND ROTATION CONTROL, WITH OR WITHOUT VARUS/VALGUS ADJUSTMENT, PREFABRICATED ITEM THAT HAS

BEEN TRIMMED, BENT, MOLDED, ASSEMBLED, OR OTHERWISE
CUSTOMIZED TO FIT A SPECIFIC PATIENT BY AN INDIVIDUAL WITH
EXPERTISE

- L1851 KNEE ORTHOSIS (KO), SINGLE UPRIGHT, THIGH AND CALF, WITH
ADJUSTABLE FLEXION AND EXTENSION JOINT (UNICENTRIC OR
POLYCENTRIC), MEDIAL-LATERAL AND ROTATION CONTROL, WITH OR
WITHOUT VARUS/VALGUS ADJUSTMENT, PREFABRICATED, OFF-THE-SHELF

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. 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. 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 about policy, claim coverage or reimbursement, please contact the DME MAC for your jurisdiction. For other questions, contact the PDAC HCPCS Helpline at the address listed above or by telephone at (877) 735-1326. The Contact Center is open Monday through Friday from 9:30 a.m. to 5:00 p.m. EST.

Sincerely,

Pricing, Data Analysis, and Coding Contract (PDAC)
Palmetto GBA, LLC
www.dmepdac.com