

September 19, 2011

COMFORTLAND MEDICAL INC
ATTN LOIS TSUI
306 OLD LARKSPUR WAY
CHAPEL HILL NC 27516

Re: Reconsideration of Coding Verification Decision

Xref #: 15809273

Product: COMFORTLAND TOUR ANKLE BRACE

Model number: CK-302

Dear Ms. Tsui:

The Pricing, Data Analysis, and Coding (PDAC) Contractor provides Healthcare Common Procedural Coding System (HCPCS) assistance to manufacturers to ensure proper coding of Durable Medical Equipment, Prosthetics, Orthotics, and Supplies (DMEPOS). The PDAC has reviewed the above listed product. It is our determination that the Medicare HCPCS code to use when billing the four Durable Medical Equipment Medicare Administrative Contractors (DME MACs) is:

L1902 - ANKLE FOOT ORTHOSIS, ANKLE GAUNTLET, PREFABRICATED,
INCLUDES FITTING AND ADJUSTMENT

The code previously assigned to this product, L1906, requires the orthosis to provide multiligamentous support. In order for an ankle orthosis to provide multiligamentous support to the ankle, it must have a rigid foot plate. This foot plate, which by means of attachment to each side of the ankle, provides functional tracking and offers hind foot and mid foot stability during ambulation. This, in conjunction with wrap around straps and the inherent gauntlet design, offer areas of multiligamentous support as described by the code. This product is an ankle gauntlet and meets the description of code L1902.

The PDAC provides coding decisions based on the coding guidelines established by the Local Coverage Determination (LCD) and associated policy article developed by the DME MACs. All products submitted to PDAC for a coding verification review are carefully examined by coders and professionals following a formal, standardized process.

From time to time questions come up with the DME MACs and PDAC as to whether a proper coding decision on a product has been made. When such a question arises, the PDAC will re-review the product and render a decision. If the decision is to change the code assigned, the manufacturer of the product is notified. The manufacturer then has the option to ask for a reconsideration, as outlined on our web site.

This coding decision will be available within ten (10) working days on the Durable Medical Equipment Coding System (DMECS), which is located on the PDAC web site, www.dmepdac.com. Please take the time to verify that this coding decision is correctly reflected in DMECS.

It is the responsibility of manufacturers and distributors to notify the PDAC immediately of any changes involving their products, related to their current listing on the Product Classification List (PCL) on DMECS. Further information for requesting updates to the PCL can be found on the PDAC website at <https://www.dmepdac.com/review/notifying.html>.

The assignment of a HCPCS code to this product is not an approval or endorsement of the product by Medicare or Noridian Administrative Services, LLC; nor does it imply or guarantee claim reimbursement or coverage. If you have questions about claim coverage or reimbursement, please contact the DME MAC for your jurisdiction.

If you disagree with this decision, you may request a reconsideration within 45 days of the date of this letter. To request a reconsideration, complete the Reconsideration Request form located on the PDAC web site at <https://www.dmepdac.com/review/requesting.html>. If your request for a reconsideration is made after the 45-day time frame, we will treat it as a coding verification review request and require a new application and documentation to support the request.

If you have questions about claim coverage or reimbursement, please contact the DME MAC for your jurisdiction. For other questions, contact the PDAC Contact Center at the address listed above or by telephone at (877) 735-1326. The Contact Center is open Monday through Friday from 8:30 a.m. to 4 p.m. CT.

PDAC
Noridian Administrative Services, LLC
www.dmepdac.com