Local Coverage Determination (LCD): Ankle-Foot/Knee-Ankle-Foot Orthosis (L33686)

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Contractor Information

CONTRACTOR NAME	CONTRACT TYPE	CONTRACT NUMBER	JURISDICTION	STATE(S)
CGS Administrators, LLC	DME MAC	17013 - DME MAC	J-B	Illinois Indiana Kentucky Michigan Minnesota Ohio Wisconsin
CGS Administrators, LLC	DME MAC	18003 - DME MAC	J-C	Alabama Arkansas Colorado Florida Georgia Louisiana Mississippi New Mexico North Carolina Oklahoma Puerto Rico South Carolina Tennessee Texas Virgin Islands Virginia West Virginia
Noridian Healthcare Solutions, LLC	DME MAC	16013 - DME MAC	J-A	Connecticut Delaware District of Columbia Maine Maryland Massachusetts New Hampshire New Jersey New York - Entire State Pennsylvania Rhode Island Vermont

CONTRACTOR NAME	CONTRACT TYPE	CONTRACT NUMBER	JURISDICTION	STATE(S)
Noridian Healthcare Solutions, LLC	DME MAC	19003 - DME MAC	J-D	Alaska American Samoa Arizona California - Entire State Guam Hawaii Idaho Iowa Kansas Missouri - Entire State Montana Nebraska Nevada North Dakota Northern Mariana Islands Oregon South Dakota Utah Washington Wyoming

LCD Information

Document Information

LCD ID
L33686

LCD Title
Ankle-Foot/Knee-Ankle-Foot Orthosis

Proposed LCD in Comment Period
N/A

Source Proposed LCD
N/A

AMA CPT / ADA CDT / AHA NUBC Copyright Statement

Original Effective Date
For services performed on or after 10/01/2015

Revision Effective Date
For services performed on or after 01/01/2020

Revision Ending Date
N/A

Retirement Date
N/A

Notice Period Start Date
N/A

Notice Period End Date
N/A

CMS National Coverage Policy

None

Coverage Guidance

Coverage Indications, Limitations, and/or Medical Necessity

For any item to be covered by Medicare, it must 1) be eligible for a defined Medicare benefit category, 2) be reasonable and necessary for the diagnosis or treatment of illness or injury or to improve the functioning of a malformed body member, and 3) meet all other applicable Medicare statutory and regulatory requirements.

The purpose of a Local Coverage Determination (LCD) is to provide information regarding "reasonable and necessary" criteria based on Social Security Act § 1862(a)(1)(A) provisions.

In addition to the "reasonable and necessary" criteria contained in this LCD there are other payment rules, which are discussed in the following documents, that must also be met prior to Medicare reimbursement:

- The LCD-related Standard Documentation Requirements Article, located at the bottom of this policy under the Related Local Coverage Documents section.
- The LCD-related Policy Article, located at the bottom of this policy under the Related Local Coverage

Documents section.
- Refer to the Supplier Manual for additional information on documentation requirements.
- Refer to the DME MAC web sites for additional bulletin articles and other publications related to this LCD.

For the items addressed in this LCD, the "reasonable and necessary" criteria, based on Social Security Act § 1862(a)(1)(A) provisions, are defined by the following coverage indications, limitations and/or medical necessity.

For Ankle-Foot Orthoses (AFO) and Knee-Ankle-Foot Orthoses (KAFO) definitions of off-the-shelf and custom fitted, refer to the CODING GUIDELINES section in the LCD-related Policy Article.

AFOs NOT USED DURING AMBULATION:

An L4396 or L4397 (Static or dynamic positioning ankle-foot orthosis) is covered if either all of criteria 1 - 4 or criterion 5 is met:

1. Plantar flexion contracture of the ankle (refer to the Group 1 Codes in the ICD-10 code list in the LCD-related Policy Article for applicable diagnoses) with dorsiflexion on passive range of motion testing of at least 10 degrees (i.e., a nonfixed contracture); and,
2. Reasonable expectation of the ability to correct the contracture; and,
3. Contracture is interfering or expected to interfere significantly with the beneficiary's functional abilities; and,
4. Used as a component of a therapy program which includes active stretching of the involved muscles and/or tendons.
5. The beneficiary has plantar fasciitis (refer to the Group 1 Codes in the ICD-10 code list in the LCD-related Policy Article for applicable diagnoses).

If an L4396 or L4397 is used for the treatment of a plantar flexion contracture, the pre-treatment passive range of motion must be measured with a goniometer and documented in the medical record. There must be documentation of an appropriate stretching program carried out by professional staff (in a nursing facility) or caregiver (at home).

An L4396 or L4397 and replacement interface (L4392) will be denied as not reasonable and necessary if the contracture is fixed. Codes L4396, L4397 and L4392 will be denied as not reasonable and necessary for a beneficiary with a foot drop but without an ankle flexion contracture. A component of a static/dynamic AFO that is used to address positioning of the knee or hip will be denied as not reasonable and necessary because the effectiveness of this type of component is not established.

If code L4396 or L4397 is covered, a replacement interface (L4392) is covered as long as the beneficiary continues to meet indications and other coverage rules for the splint. Coverage of a replacement interface is limited to a maximum of one (1) per 6 months. Additional interfaces will be denied as not reasonable and necessary.

Medicare does not reimburse for a foot drop splint/recumbent positioning device (L4398) or replacement interface (L4394). A foot drop splint/recumbent positioning device and replacement interface will be denied as not reasonable and necessary in a beneficiary with foot drop who is nonambulatory because there are other more appropriate treatment modalities.

AFOs AND KAFOs USED DURING AMBULATION:

Ankle-foot orthoses (AFO) described by codes L1900, L1902, L1904, L1906, L1907, L1910, L1920, L1930, L1932, L1940, L1945, L1950, L1951, L1960, L1970, L1971, L1980, L1990, L2106, L2108, L2112, L2114, L2116, L4350, L4360, L4361, L4386, L4387 and L4631 are covered for ambulatory beneficiaries with weakness or deformity of the foot and ankle, who:

1. Require stabilization for medical reasons, and,

2. Have the potential to benefit functionally.

Knee-ankle-foot orthoses (KAFO) described by codes L2000, L2005, L2010, L2020, L2030, L2034, L2035, L2036, L2037, L2038, L2126, L2128, L2132, L2134, L2136, and L4370 are covered for ambulatory beneficiaries for whom an ankle-foot orthosis is covered and for whom additional knee stability is required.

If the basic coverage criteria for an AFO or KAFO are not met, the orthosis will be denied as not reasonable and necessary.

AFOs and KAFOs that are custom-fabricated are covered for ambulatory beneficiaries when the basic coverage criteria listed above and one of the following criteria are met:

1. The beneficiary could not be fit with a prefabricated AFO; or,
2. The condition necessitating the orthosis is expected to be permanent or of longstanding duration (more than 6 months); or,
3. There is a need to control the knee, ankle or foot in more than one plane; or,
4. The beneficiary has a documented neurological, circulatory, or orthopedic status that requires custom fabricating over a model to prevent tissue injury; or,
5. The beneficiary has a healing fracture which lacks normal anatomical integrity or anthropometric proportions.

If a custom fabricated orthosis is provided but basic coverage criteria above and the additional criteria 1-5 for a custom fabricated orthosis are not met, the custom fabricated orthosis will be denied as not reasonable and necessary.

L coded additions to AFOs and KAFOs (L2180, L2182, L2184, L2186, L2188, L2190, L2192, L2200, L2210, L2220, L2230, L2232, L2240, L2250, L2260, L2265, L2270, L2275, L2280, L2300, L2310, L2320, L2330, L2335, L2340, L2350, L2360, L2370, L2375, L2380, L2385, L2387, L2390, L2395, L2397, L2405, L2415, L2425, L2430, L2492, L2500, L2510, L2520, L2525, L2526, L2530, L2540, L2550, L2750, L2755, L2760, L2768, L2780, L2785, L2795, L2800, L2810, L2820, L2830) will be denied as not reasonable and necessary if either the base orthosis is not reasonable and necessary or the specific addition is not reasonable and necessary.

Concentric adjustable torsion style mechanisms used to assist knee joint extension are coded as L2999 and are covered for beneficiaries who require knee extension assist in the absence of any co-existing joint contracture.

Concentric adjustable torsion style mechanisms used to assist ankle joint plantarflexion or dorsiflexion are coded as L2999 and are covered for beneficiaries who require ankle plantar or dorsiflexion assist in the absence of any co-existing joint contracture.

Concentric adjustable torsion style mechanisms used for the treatment of contractures, regardless of any co-existing condition(s), are coded as E1810 and/or E1815 and are covered under the Durable Medical Equipment benefit (refer to the CODING GUIDELINES section in the LCD-related Policy Article).

Claims for devices incorporating concentric adjustable torsion style mechanisms used for the treatment of any joint contracture and coded as L2999 will be denied as incorrect coding.

Refer to the Orthopedic Footwear policy for information on coverage of shoes and related items which are an integral part of a brace.

Replacement components (e.g., soft interfaces) that are provided on a routine basis, without regard to whether the

original item is worn out, are covered under the refill requirements.

GENERAL

A Standard Written Order (SWO) must be communicated to the supplier before a claim is submitted. If the supplier bills for an item addressed in this policy without first receiving a completed SWO, the claim shall be denied as not reasonable and necessary.

For Durable Medical Equipment, Prosthetics, Orthotics and Supplies (DMEPOS) base items that require a Written Order Prior to Delivery (WOPD), the supplier must have received a signed SWO before the DMEPOS item is delivered to a beneficiary. If a supplier delivers a DMEPOS item without first receiving a WOPD, the claim shall be denied as not reasonable and necessary. Refer to the LCD-related Policy Article, located at the bottom of this policy under the Related Local Coverage Documents section.

For DMEPOS base items that require a WOPD, and also require separately billed associated options, accessories, and/or supplies, the supplier must have received a WOPD which lists the base item and which may list all the associated options, accessories, and/or supplies that are separately billed prior to the delivery of the items. In this scenario, if the supplier separately bills for associated options, accessories, and/or supplies without first receiving a completed and signed WOPD of the base item prior to delivery, the claim(s) shall be denied as not reasonable and necessary.

An item/service is correctly coded when it meets all the coding guidelines listed in CMS HCPCS guidelines, LCDs, LCD-related Policy Articles, or DME MAC articles. Claims that do not meet coding guidelines shall be denied as not reasonable and necessary/incorrectly coded.

Proof of delivery (POD) is a Supplier Standard and DMEPOS suppliers are required to maintain POD documentation in their files. Proof of delivery documentation must be made available to the Medicare contractor upon request. All services that do not have appropriate proof of delivery from the supplier shall be denied as not reasonable and necessary.

Summary of Evidence

N/A

Analysis of Evidence
(Rationale for Determination)

N/A

Coding Information

CPT/HCPCS Codes

Group 1 Paragraph:

The appearance of a code in this section does not necessarily indicate coverage.

HCPCS MODIFIERS:

EY - No physician or other licensed health care provider order for this item or service

GA – Waiver of liability statement issued as required by payer policy, individual case

GZ – Item or service expected to be denied as not reasonable and necessary

KX - Requirements specified in the medical policy have been met

LT - Left Side

RT - Right Side

HCPCS CODES:

Group 1 Codes:

CODE	DESCRIPTION
A4467	BELT, STRAP, SLEEVE, GARMENT, OR COVERING, ANY TYPE
A9283	FOOT PRESSURE OFF LOADING/SUPPORTIVE DEVICE, ANY TYPE, EACH
A9285	INVERSION/EVERSION CORRECTION DEVICE
L1900	ANKLE FOOT ORTHOSIS, SPRING WIRE, DORSIFLEXION ASSIST CALF BAND, CUSTOM FABRICATED
L1902	ANKLE ORTHOSIS, ANKLE GAUNTLET OR SIMILAR, WITH OR WITHOUT JOINTS, PREFABRICATED, OFF-THE-SHELF
L1904	ANKLE ORTHOSIS, ANKLE GAUNTLET OR SIMILAR, WITH OR WITHOUT JOINTS, CUSTOM FABRICATED
L1906	ANKLE FOOT ORTHOSIS, MULTILIGAMENTOUS ANKLE SUPPORT, PREFABRICATED, OFF-THE-SHELF

CODE	DESCRIPTION
L1907	ANKLE ORTHOSIS, SUPRAMALLEOLAR WITH STRAPS, WITH OR WITHOUT INTERFACE/PADS, CUSTOM FABRICATED
L1910	ANKLE FOOT ORTHOSIS, POSTERIOR, SINGLE BAR, CLASP ATTACHMENT TO SHOE COUNTER, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L1920	ANKLE FOOT ORTHOSIS, SINGLE UPRIGHT WITH STATIC OR ADJUSTABLE STOP (PHELPS OR PERLSTEIN TYPE), CUSTOM FABRICATED
L1930	ANKLE FOOT ORTHOSIS, PLASTIC OR OTHER MATERIAL, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L1932	AFO, RIGID ANTERIOR TIBIAL SECTION, TOTAL CARBON FIBER OR EQUAL MATERIAL, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L1940	ANKLE FOOT ORTHOSIS, PLASTIC OR OTHER MATERIAL, CUSTOM FABRICATED
L1945	ANKLE FOOT ORTHOSIS, PLASTIC, RIGID ANTERIOR TIBIAL SECTION (FLOOR REACTION), CUSTOM FABRICATED
L1950	ANKLE FOOT ORTHOSIS, SPIRAL, (INSTITUTE OF REHABILITATIVE MEDICINE TYPE), PLASTIC, CUSTOM FABRICATED
L1951	ANKLE FOOT ORTHOSIS, SPIRAL, (INSTITUTE OF REHABILITATIVE MEDICINE TYPE), PLASTIC OR OTHER MATERIAL, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L1960	ANKLE FOOT ORTHOSIS, POSTERIOR SOLID ANKLE, PLASTIC, CUSTOM FABRICATED
L1970	ANKLE FOOT ORTHOSIS, PLASTIC WITH ANKLE JOINT, CUSTOM FABRICATED
L1971	ANKLE FOOT ORTHOSIS, PLASTIC OR OTHER MATERIAL WITH ANKLE JOINT, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L1980	ANKLE FOOT ORTHOSIS, SINGLE UPRIGHT FREE PLANTAR DORSIFLEXION, SOLID STIRRUP, CALF BAND/CUFF (SINGLE BAR 'BK' ORTHOSIS), CUSTOM FABRICATED
L1990	ANKLE FOOT ORTHOSIS, DOUBLE UPRIGHT FREE PLANTAR DORSIFLEXION, SOLID STIRRUP, CALF BAND/CUFF (DOUBLE BAR 'BK' ORTHOSIS), CUSTOM FABRICATED
L2000	KNEE ANKLE FOOT ORTHOSIS, SINGLE UPRIGHT, FREE KNEE, FREE ANKLE, SOLID STIRRUP, THIGH AND CALF BANDS/CUFFS (SINGLE BAR 'AK' ORTHOSIS), CUSTOM FABRICATED
L2005	KNEE ANKLE FOOT ORTHOSIS, ANY MATERIAL, SINGLE OR DOUBLE UPRIGHT, STANCE CONTROL, AUTOMATIC LOCK AND SWING PHASE RELEASE, ANY TYPE ACTIVATION, INCLUDES ANKLE JOINT, ANY TYPE, CUSTOM FABRICATED
L2006	KNEE ANKLE FOOT DEVICE, ANY MATERIAL, SINGLE OR DOUBLE UPRIGHT, SWING AND STANCE PHASE MICROPROCESSOR CONTROL WITH ADJUSTABILITY, INCLUDES ALL COMPONENTS (E.G., SENSORS, BATTERIES, CHARGER), ANY TYPE ACTIVATION, WITH OR WITHOUT ANKLE JOINT(S), CUSTOM FABRICATED
L2010	KNEE ANKLE FOOT ORTHOSIS, SINGLE UPRIGHT, FREE ANKLE, SOLID STIRRUP,

CODE	DESCRIPTION
	THIGH AND CALF BANDS/CUFFS (SINGLE BAR 'AK' ORTHOSIS), WITHOUT KNEE JOINT, CUSTOM FABRICATED
L2020	KNEE ANKLE FOOT ORTHOSIS, DOUBLE UPRIGHT, FREE ANKLE, SOLID STIRRUP, THIGH AND CALF BANDS/CUFFS (DOUBLE BAR 'AK' ORTHOSIS), CUSTOM FABRICATED
L2030	KNEE ANKLE FOOT ORTHOSIS, DOUBLE UPRIGHT, FREE ANKLE, SOLID STIRRUP, THIGH AND CALF BANDS/CUFFS, (DOUBLE BAR 'AK' ORTHOSIS), WITHOUT KNEE JOINT, CUSTOM FABRICATED
L2034	KNEE ANKLE FOOT ORTHOSIS, FULL PLASTIC, SINGLE UPRIGHT, WITH OR WITHOUT FREE MOTION KNEE, MEDIAL LATERAL ROTATION CONTROL, WITH OR WITHOUT FREE MOTION ANKLE, CUSTOM FABRICATED
L2035	KNEE ANKLE FOOT ORTHOSIS, FULL PLASTIC, STATIC (PEDIATRIC SIZE), WITHOUT FREE MOTION ANKLE, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L2036	KNEE ANKLE FOOT ORTHOSIS, FULL PLASTIC, DOUBLE UPRIGHT, WITH OR WITHOUT FREE MOTION KNEE, WITH OR WITHOUT FREE MOTION ANKLE, CUSTOM FABRICATED
L2037	KNEE ANKLE FOOT ORTHOSIS, FULL PLASTIC, SINGLE UPRIGHT, WITH OR WITHOUT FREE MOTION KNEE, WITH OR WITHOUT FREE MOTION ANKLE, CUSTOM FABRICATED
L2038	KNEE ANKLE FOOT ORTHOSIS, FULL PLASTIC, WITH OR WITHOUT FREE MOTION KNEE, MULTI-AXIS ANKLE, CUSTOM FABRICATED
L2106	ANKLE FOOT ORTHOSIS, FRACTURE ORTHOSIS, TIBIAL FRACTURE CAST ORTHOSIS, THERMOPLASTIC TYPE CASTING MATERIAL, CUSTOM FABRICATED
L2108	ANKLE FOOT ORTHOSIS, FRACTURE ORTHOSIS, TIBIAL FRACTURE CAST ORTHOSIS, CUSTOM FABRICATED
L2112	ANKLE FOOT ORTHOSIS, FRACTURE ORTHOSIS, TIBIAL FRACTURE ORTHOSIS, SOFT, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L2114	ANKLE FOOT ORTHOSIS, FRACTURE ORTHOSIS, TIBIAL FRACTURE ORTHOSIS, SEMI-RIGID, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L2116	ANKLE FOOT ORTHOSIS, FRACTURE ORTHOSIS, TIBIAL FRACTURE ORTHOSIS, RIGID, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L2126	KNEE ANKLE FOOT ORTHOSIS, FRACTURE ORTHOSIS, FEMORAL FRACTURE CAST ORTHOSIS, THERMOPLASTIC TYPE CASTING MATERIAL, CUSTOM FABRICATED
L2128	KNEE ANKLE FOOT ORTHOSIS, FRACTURE ORTHOSIS, FEMORAL FRACTURE CAST ORTHOSIS, CUSTOM FABRICATED
L2132	KAFO, FRACTURE ORTHOSIS, FEMORAL FRACTURE CAST ORTHOSIS, SOFT, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L2134	KAFO, FRACTURE ORTHOSIS, FEMORAL FRACTURE CAST ORTHOSIS, SEMI-RIGID,

CODE	DESCRIPTION
	PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L2136	KAFO, FRACTURE ORTHOSIS, FEMORAL FRACTURE CAST ORTHOSIS, RIGID, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT
L2180	ADDITION TO LOWER EXTREMITY FRACTURE ORTHOSIS, PLASTIC SHOE INSERT WITH ANKLE JOINTS
L2182	ADDITION TO LOWER EXTREMITY FRACTURE ORTHOSIS, DROP LOCK KNEE JOINT
L2184	ADDITION TO LOWER EXTREMITY FRACTURE ORTHOSIS, LIMITED MOTION KNEE JOINT
L2186	ADDITION TO LOWER EXTREMITY FRACTURE ORTHOSIS, ADJUSTABLE MOTION KNEE JOINT, LERMAN TYPE
L2188	ADDITION TO LOWER EXTREMITY FRACTURE ORTHOSIS, QUADRILATERAL BRIM
L2190	ADDITION TO LOWER EXTREMITY FRACTURE ORTHOSIS, WAIST BELT
L2192	ADDITION TO LOWER EXTREMITY FRACTURE ORTHOSIS, HIP JOINT, PELVIC BAND, THIGH FLANGE, AND PELVIC BELT
L2200	ADDITION TO LOWER EXTREMITY, LIMITED ANKLE MOTION, EACH JOINT
L2210	ADDITION TO LOWER EXTREMITY, DORSIFLEXION ASSIST (PLANTAR FLEXION RESIST), EACH JOINT
L2220	ADDITION TO LOWER EXTREMITY, DORSIFLEXION AND PLANTAR FLEXION ASSIST/RESIST, EACH JOINT
L2230	ADDITION TO LOWER EXTREMITY, SPLIT FLAT CALIPER STIRRUPS AND PLATE ATTACHMENT
L2232	ADDITION TO LOWER EXTREMITY ORTHOSIS, ROCKER BOTTOM FOR TOTAL CONTACT ANKLE FOOT ORTHOSIS, FOR CUSTOM FABRICATED ORTHOSIS ONLY
L2240	ADDITION TO LOWER EXTREMITY, ROUND CALIPER AND PLATE ATTACHMENT
L2250	ADDITION TO LOWER EXTREMITY, FOOT PLATE, MOLDED TO PATIENT MODEL, STIRRUP ATTACHMENT
L2260	ADDITION TO LOWER EXTREMITY, REINFORCED SOLID STIRRUP (SCOTT-CRAIG TYPE)
L2265	ADDITION TO LOWER EXTREMITY, LONG TONGUE STIRRUP
L2270	ADDITION TO LOWER EXTREMITY, VARUS/VALGUS CORRECTION ('T') STRAP, PADDED/LINED OR MALLEOLUS PAD
L2275	ADDITION TO LOWER EXTREMITY, VARUS/VALGUS CORRECTION, PLASTIC MODIFICATION, PADDED/LINED
L2280	ADDITION TO LOWER EXTREMITY, MOLDED INNER BOOT
L2300	ADDITION TO LOWER EXTREMITY, ABDUCTION BAR (BILATERAL HIP INVOLVEMENT), JOINTED, ADJUSTABLE

CODE	DESCRIPTION
L2310	ADDITION TO LOWER EXTREMITY, ABDUCTION BAR-STRAIGHT
L2320	ADDITION TO LOWER EXTREMITY, NON-MOLDED LACER, FOR CUSTOM FABRICATED ORTHOSIS ONLY
L2330	ADDITION TO LOWER EXTREMITY, LACER MOLDED TO PATIENT MODEL, FOR CUSTOM FABRICATED ORTHOSIS ONLY
L2335	ADDITION TO LOWER EXTREMITY, ANTERIOR SWING BAND
L2340	ADDITION TO LOWER EXTREMITY, PRE-TIBIAL SHELL, MOLDED TO PATIENT MODEL
L2350	ADDITION TO LOWER EXTREMITY, PROSTHETIC TYPE, (BK) SOCKET, MOLDED TO PATIENT MODEL, (USED FOR 'PTB' 'AFO' ORTHOSES)
L2360	ADDITION TO LOWER EXTREMITY, EXTENDED STEEL SHANK
L2370	ADDITION TO LOWER EXTREMITY, PATTEN BOTTOM
L2375	ADDITION TO LOWER EXTREMITY, TORSION CONTROL, ANKLE JOINT AND HALF SOLID STIRRUP
L2380	ADDITION TO LOWER EXTREMITY, TORSION CONTROL, STRAIGHT KNEE JOINT, EACH JOINT
L2385	ADDITION TO LOWER EXTREMITY, STRAIGHT KNEE JOINT, HEAVY DUTY, EACH JOINT
L2387	ADDITION TO LOWER EXTREMITY, POLYCENTRIC KNEE JOINT, FOR CUSTOM FABRICATED KNEE ANKLE FOOT ORTHOSIS, EACH JOINT
L2390	ADDITION TO LOWER EXTREMITY, OFFSET KNEE JOINT, EACH JOINT
L2395	ADDITION TO LOWER EXTREMITY, OFFSET KNEE JOINT, HEAVY DUTY, EACH JOINT
L2397	ADDITION TO LOWER EXTREMITY ORTHOSIS, SUSPENSION SLEEVE
L2405	ADDITION TO KNEE JOINT, DROP LOCK, EACH
L2415	ADDITION TO KNEE LOCK WITH INTEGRATED RELEASE MECHANISM (BAIL, CABLE, OR EQUAL), ANY MATERIAL, EACH JOINT
L2425	ADDITION TO KNEE JOINT, DISC OR DIAL LOCK FOR ADJUSTABLE KNEE FLEXION, EACH JOINT
L2430	ADDITION TO KNEE JOINT, RATCHET LOCK FOR ACTIVE AND PROGRESSIVE KNEE EXTENSION, EACH JOINT
L2492	ADDITION TO KNEE JOINT, LIFT LOOP FOR DROP LOCK RING
L2500	ADDITION TO LOWER EXTREMITY, THIGH/WEIGHT BEARING, GLUTEAL/ ISCHIAL WEIGHT BEARING, RING
L2510	ADDITION TO LOWER EXTREMITY, THIGH/WEIGHT BEARING, QUADRI- LATERAL BRIM, MOLDED TO PATIENT MODEL
L2520	ADDITION TO LOWER EXTREMITY, THIGH/WEIGHT BEARING, QUADRI- LATERAL

CODE	DESCRIPTION
	BRIM, CUSTOM FITTED
L2525	ADDITION TO LOWER EXTREMITY, THIGH/WEIGHT BEARING, ISCHIAL CONTAINMENT/NARROW M-L BRIM MOLDED TO PATIENT MODEL
L2526	ADDITION TO LOWER EXTREMITY, THIGH/WEIGHT BEARING, ISCHIAL CONTAINMENT/NARROW M-L BRIM, CUSTOM FITTED
L2530	ADDITION TO LOWER EXTREMITY, THIGH-WEIGHT BEARING, LACER, NON-MOLDED
L2540	ADDITION TO LOWER EXTREMITY, THIGH/WEIGHT BEARING, LACER, MOLDED TO PATIENT MODEL
L2550	ADDITION TO LOWER EXTREMITY, THIGH/WEIGHT BEARING, HIGH ROLL CUFF
L2750	ADDITION TO LOWER EXTREMITY ORTHOSIS, PLATING CHROME OR NICKEL, PER BAR
L2755	ADDITION TO LOWER EXTREMITY ORTHOSIS, HIGH STRENGTH, LIGHTWEIGHT MATERIAL, ALL HYBRID LAMINATION/PREPREG COMPOSITE, PER SEGMENT, FOR CUSTOM FABRICATED ORTHOSIS ONLY
L2760	ADDITION TO LOWER EXTREMITY ORTHOSIS, EXTENSION, PER EXTENSION, PER BAR (FOR LINEAL ADJUSTMENT FOR GROWTH)
L2768	ORTHOTIC SIDE BAR DISCONNECT DEVICE, PER BAR
L2780	ADDITION TO LOWER EXTREMITY ORTHOSIS, NON-CORROSIVE FINISH, PER BAR
L2785	ADDITION TO LOWER EXTREMITY ORTHOSIS, DROP LOCK RETAINER, EACH
L2795	ADDITION TO LOWER EXTREMITY ORTHOSIS, KNEE CONTROL, FULL KNEECAP
L2800	ADDITION TO LOWER EXTREMITY ORTHOSIS, KNEE CONTROL, KNEE CAP, MEDIAL OR LATERAL PULL, FOR USE WITH CUSTOM FABRICATED ORTHOSIS ONLY
L2810	ADDITION TO LOWER EXTREMITY ORTHOSIS, KNEE CONTROL, CONDYLAR PAD
L2820	ADDITION TO LOWER EXTREMITY ORTHOSIS, SOFT INTERFACE FOR MOLDED PLASTIC, BELOW KNEE SECTION
L2830	ADDITION TO LOWER EXTREMITY ORTHOSIS, SOFT INTERFACE FOR MOLDED PLASTIC, ABOVE KNEE SECTION
CODE	DESCRIPTION
L2840	ADDITION TO LOWER EXTREMITY ORTHOSIS, TIBIAL LENGTH SOCK, FRACTURE OR EQUAL, EACH
L2850	ADDITION TO LOWER EXTREMITY ORTHOSIS, FEMORAL LENGTH SOCK, FRACTURE OR EQUAL, EACH
L2999	LOWER EXTREMITY ORTHOSES, NOT OTHERWISE SPECIFIED
L4002	REPLACEMENT STRAP, ANY ORTHOSIS, INCLUDES ALL COMPONENTS, ANY LENGTH, ANY TYPE

CODE	DESCRIPTION
L4010	REPLACE TRILATERAL SOCKET BRIM
L4020	REPLACE QUADRILATERAL SOCKET BRIM, MOLDED TO PATIENT MODEL
L4030	REPLACE QUADRILATERAL SOCKET BRIM, CUSTOM FITTED
L4040	REPLACE MOLDED THIGH LACER, FOR CUSTOM FABRICATED ORTHOSIS ONLY
L4045	REPLACE NON-MOLDED THIGH LACER, FOR CUSTOM FABRICATED ORTHOSIS ONLY
L4050	REPLACE MOLDED CALF LACER, FOR CUSTOM FABRICATED ORTHOSIS ONLY
L4055	REPLACE NON-MOLDED CALF LACER, FOR CUSTOM FABRICATED ORTHOSIS ONLY
L4060	REPLACE HIGH ROLL CUFF
L4070	REPLACE PROXIMAL AND DISTAL UPRIGHT FOR KAFO
L4080	REPLACE METAL BANDS KAFO, PROXIMAL THIGH
L4090	REPLACE METAL BANDS KAFO-AFO, CALF OR DISTAL THIGH
L4100	REPLACE LEATHER CUFF KAFO, PROXIMAL THIGH
L4110	REPLACE LEATHER CUFF KAFO-AFO, CALF OR DISTAL THIGH
L4130	REPLACE PRETIBIAL SHELL
L4205	REPAIR OF ORTHOTIC DEVICE, LABOR COMPONENT, PER 15 MINUTES
L4210	REPAIR OF ORTHOTIC DEVICE, REPAIR OR REPLACE MINOR PARTS
L4350	ANKLE CONTROL ORTHOSIS, STIRRUP STYLE, RIGID, INCLUDES ANY TYPE INTERFACE (E.G., PNEUMATIC, GEL), PREFABRICATED, OFF-THE-SHELF
L4360	WALKING BOOT, PNEUMATIC AND/OR VACUUM, WITH OR WITHOUT JOINTS, WITH OR WITHOUT INTERFACE MATERIAL, PREFABRICATED ITEM THAT HAS BEEN TRIMMED, BENT, MOLDED, ASSEMBLED, OR OTHERWISE CUSTOMIZED TO FIT A SPECIFIC PATIENT BY AN INDIVIDUAL WITH EXPERTISE
L4361	WALKING BOOT, PNEUMATIC AND/OR VACUUM, WITH OR WITHOUT JOINTS, WITH OR WITHOUT INTERFACE MATERIAL, PREFABRICATED, OFF-THE-SHELF
L4370	PNEUMATIC FULL LEG SPLINT, PREFABRICATED, OFF-THE-SHELF
L4386	WALKING BOOT, NON-PNEUMATIC, WITH OR WITHOUT JOINTS, WITH OR WITHOUT INTERFACE MATERIAL, PREFABRICATED ITEM THAT HAS BEEN TRIMMED, BENT, MOLDED, ASSEMBLED, OR OTHERWISE CUSTOMIZED TO FIT A SPECIFIC PATIENT BY AN INDIVIDUAL WITH EXPERTISE
L4387	WALKING BOOT, NON-PNEUMATIC, WITH OR WITHOUT JOINTS, WITH OR WITHOUT INTERFACE MATERIAL, PREFABRICATED, OFF-THE-SHELF
L4392	REPLACEMENT, SOFT INTERFACE MATERIAL, STATIC AFO
L4394	REPLACE SOFT INTERFACE MATERIAL, FOOT DROP SPLINT
L4396	STATIC OR DYNAMIC ANKLE FOOT ORTHOSIS, INCLUDING SOFT INTERFACE MATERIAL, ADJUSTABLE FOR FIT, FOR POSITIONING, MAY BE USED FOR MINIMAL

CODE	DESCRIPTION
	AMBULATION, PREFABRICATED ITEM THAT HAS BEEN TRIMMED, BENT, MOLDED, ASSEMBLED, OR OTHERWISE CUSTOMIZED TO FIT A SPECIFIC PATIENT BY AN INDIVIDUAL WITH EXPERTISE
L4397	STATIC OR DYNAMIC ANKLE FOOT ORTHOSIS, INCLUDING SOFT INTERFACE MATERIAL, ADJUSTABLE FOR FIT, FOR POSITIONING, MAY BE USED FOR MINIMAL AMBULATION, PREFABRICATED, OFF-THE-SHELF
L4398	FOOT DROP SPLINT, RECUMBENT POSITIONING DEVICE, PREFABRICATED, OFF-THE-SHELF
L4631	ANKLE FOOT ORTHOSIS, WALKING BOOT TYPE, VARUS/VALGUS CORRECTION, ROCKER BOTTOM, ANTERIOR TIBIAL SHELL, SOFT INTERFACE, CUSTOM ARCH SUPPORT, PLASTIC OR OTHER MATERIAL, INCLUDES STRAPS AND CLOSURES, CUSTOM FABRICATED

General Information

Associated Information

DOCUMENTATION REQUIREMENTS
Section 1833(e) of the Social Security Act precludes payment to any provider of services unless "there has been furnished such information as may be necessary in order to determine the amounts due such provider.” It is expected that the beneficiary's medical records will reflect the need for the care provided. The beneficiary's medical records include the treating practitioner's office records, hospital records, nursing home records, home health agency records, records from other healthcare professionals and test reports. This documentation must be available upon request.

GENERAL DOCUMENTATION REQUIREMENTS

In order to justify payment for DMEPOS items, suppliers must meet the following requirements:

- SWO
- Medical Record Information (including continued need/use if applicable)
- Correct Coding
- Proof of Delivery

Refer to the LCD-related Standard Documentation Requirements article, located at the bottom of this policy under the Related Local Coverage Documents section for additional information regarding these requirements.

Refer to the Supplier Manual for additional information on documentation requirements.

Refer to the DME MAC web sites for additional bulletin articles and other publications related to this LCD.

POLICY SPECIFIC DOCUMENTATION REQUIREMENTS

Items covered in this LCD have additional policy-specific requirements that must be met prior to Medicare reimbursement.

Refer to the LCD-related Policy article, located at the bottom of this policy under the Related Local Coverage Documents section for additional information.

Miscellaneous

Appendices

Utilization Guidelines

Refer to Coverage Indications, Limitations and/or Medical Necessity

Sources of Information

N/A

Bibliography

N/A

Revision History Information

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION	REASON(S) FOR CHANGE
01/01/2020	R8	Revision Effective Date: 01/01/2020 COVERAGE INDICATIONS, LIMITATIONS, AND/OR MEDICAL NECESSITY: Revised: Format of HCPCS code references, from code 'spans' to individually-listed HCPCS Revised: Order information as a result of Final Rule 1713 HCPCS CODES: Revised: HCPCS L2006 code description per quarterly HCPCS code update CODING INFORMATION: Removed: Field titled "Bill Type" Removed: Field titled "Revenue Codes" Removed: Field titled "ICD-10 Codes that Support Medical Necessity" Removed: Field titled "ICD-10 Codes that DO NOT Support Medical Necessity" Removed: Field titled "Additional ICD-10 Information" DOCUMENTATION REQUIREMENTS: Revised: "physician's" to "treating practitioner's" GENERAL DOCUMENTATION REQUIREMENTS: Revised: Prescriptions (orders) to SWO	• Provider Education/Guidance • Other

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION	REASON(S) FOR CHANGE
		02/20/2020: Pursuant to the 21st Century Cures Act, these revisions do not require notice and comment because they are due to non-discretionary coverage updates reflective of CMS FR-1713, HCPCS code changes, and non-substantive corrections (listing individual HCPCS codes instead of a HCPCS code-span). During the exercise of listing individual HCPCS codes, L2006 had been inadvertently added because it fell within a code span and is being deleted.	
01/01/2020	R7	Revision Effective Date: 01/01/2020 COVERAGE INDICATIONS, LIMITATIONS, AND/OR MEDICAL NECESSITY: Removed: Statement to refer to ICD-10 Codes that are Covered section in the LCD-related PA Added: Statement to refer to ICD-10 code list in the LCD-related Policy Article HCPCS CODES: Added: HCPCS L2006 to Group 1 codes, per annual HCPCS code release	• Revisions Due To CPT/HCPCS Code Changes
01/01/2019	R6	Revision Effective Date: 01/01/2019 COVERAGE INDICATIONS, LIMITATIONS, AND/OR MEDICAL NECESSITY: Removed: Statement to refer to diagnosis code section below Added: Refer to Covered ICD-10 Codes in the LCD-related Policy Article ICD-10 CODES THAT SUPPORT MEDICAL NECESSITY: Moved: All diagnosis codes to the LCD-related Policy Article diagnosis code section per CMS instruction ICD-10 CODES THAT DO NOT SUPPORT MEDICAL NECESSITY: Moved: Statement about noncovered diagnosis codes moved to LCD-related Policy Article noncovered diagnosis code section per CMS instruction	• Other (ICD-10 code relocation per CMS instruction)
01/01/2017	R5	No changes have been made to this LCD *03/29/2018: At this time 21st Century Cures Act will apply to new and revised LCDs that restrict coverage which requires comment and notice. This revision is not a restriction to the*	• Other

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION	REASON(S) FOR CHANGE
		coverage determination; and, therefore not all the fields included on the LCD are applicable as noted in this policy.	
01/01/2017	R4	Revision Effective Date: 01/01/2017: COVERAGE INDICATIONS, LIMITATIONS AND/OR MEDICAL NECESSITY: Removed: Standard Documentation Language Added: New reference language and Directions to Standard Documentation Requirements Added: General Requirements HCPCS CODES: Added: HCPCS Code A4467 & A9285 Deleted: HCPCS Code A4466 Revised: HCPCS Code L1906 ICD-10 CODES THAT SUPPORT MEDICAL NECESSITY: Deleted: ICD-10 Diagnoses (M14.661, M14.662, M14.669) for L4631; diagnoses not pertinent to this orthosis DOCUMENTATION REQUIREMENTS: Removed: Standard Documentation Language Added: General Documentation Requirements Added: New reference language and Directions to Standard Documentation Requirements POLICY SPECIFIC DOCUMENTATION REQUIREMENTS: Removed: Standard Documentation Language Added: Directions to Standard Documentation Requirements Removed: Information under Miscellaneous and Appendices RELATED LOCAL COVERAGE DOCUMENTS: Added: LCD-related Standard Documentation Requirements article	• Provider Education/Guidance • Revisions Due To ICD-10-CM Code Changes • Revisions Due To CPT/HCPCS Code Changes
07/01/2016	R3	Effective July 1, 2016 oversight for DME MAC LCDs is the responsibility of CGS Administrators, LLC 18003 and 17013 and Noridian Healthcare Solutions, LLC 19003 and 16013. No other changes have been made to the LCDs.	• Change in Assigned States or Affiliated Contract Numbers
01/01/2016	R2	**Revision Effective Date: 01/01/2016:** COVERAGE INDICATIONS, LIMITATIONS AND/OR MEDICAL NECESSITY:	• Provider Education/Guidance • Revisions Due To

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION	REASON(S) FOR CHANGE
		Added: L4361 "clerical correction" HCPCS CODES: Revised: L1902 and L1904 long narrative description DOCUMENTATION REQUIREMENTS: Revised: Standard Documentation Language to remove start date verbiage from Prescription Requirements (Effective 11/5/2015) Moved: Repair/Replacement verbiage to correct location Updated: Miscellaneous section when billing L2999	CPT/HCPCS Code Changes
10/01/2015	R1	Revision Effective Date: 05/01/2015 COVERAGE INDICATIONS, LIMITATIONS AND/OR MEDICAL NECESSITY: Revised: Standard Documentation Language to add covered prior to a beneficiary's Medicare eligibility DOCUMENTATION REQUIREMENTS: Added: Continued Need & Continue Use Revised: Standard Documentation Language to add who can enter date of delivery date on the POD Added: Instructions for Equipment Retained from a Prior Payer POLICY SPECIFIC DOCUMENTATION REQUIREMENTS: Updated: Documentation responsibilities for prefabricated vs. custom fabricated devices to reflect revision of April 2015 bulletin article Revised: Repair to beneficiary-owned DMEPOS Revised: Instructions for HCPCS L2999	• Provider Education/Guidance

Associated Documents

Attachments

N/A

Related Local Coverage Documents

Article(s)
A52457 - Ankle-Foot/Knee-Ankle-Foot Orthoses - Policy Article
A55426 - Standard Documentation Requirements for All Claims Submitted to DME MACs

Related National Coverage Documents

N/A

Public Version(s)

Updated on 02/14/2020 with effective dates 01/01/2020 - N/A
Updated on 12/13/2019 with effective dates 01/01/2020 - N/A

Updated on 03/22/2019 with effective dates 01/01/2019 - 12/31/2019

Some older versions have been archived. Please visit the MCD Archive Site to retrieve them.

Keywords

N/A

END OF LOCAL COVERAGE DETERMINATION

Per the Code of Federal Regulations, 42 C.F.R § 426. 325, only those portions of the currently effective Local Coverage Determination (LCD) that are based on section 1862(a)(1)(A) of the Social Security Act, may be challenged through an acceptable complaint as described in 42 C.F.R § 426.400. Also, per 42 C.F.R § 426.325 items that are not reviewable, and therefore cannot be challenged, include the Policy Article. Please note the distinction of the documents when reviewing the materials.

Local Coverage Article:
Ankle-Foot/Knee-Ankle-Foot Orthoses - Policy Article (A52457)

Links in PDF documents are not guaranteed to work. To follow a web link, please use the MCD Website.

Contractor Information

CONTRACTOR NAME	CONTRACT TYPE	CONTRACT NUMBER	JURISDICTION	STATE(S)
CGS Administrators, LLC	DME MAC	17013 - DME MAC	J-B	Illinois Indiana Kentucky Michigan Minnesota Ohio Wisconsin
CGS Administrators, LLC	DME MAC	18003 - DME MAC	J-C	Alabama Arkansas Colorado Florida Georgia Louisiana Mississippi New Mexico North Carolina Oklahoma Puerto Rico South Carolina Tennessee Texas Virgin Islands Virginia West Virginia
Noridian Healthcare Solutions, LLC	DME MAC	16013 - DME MAC	J-A	Connecticut Delaware District of Columbia Maine Maryland Massachusetts New Hampshire New Jersey New York - Entire State Pennsylvania Rhode Island Vermont

CONTRACTOR NAME	CONTRACT TYPE	CONTRACT NUMBER	JURISDICTION	STATE(S)
Noridian Healthcare Solutions, LLC	DME MAC	19003 - DME MAC	J-D	Alaska American Samoa Arizona California - Entire State Guam Hawaii Idaho Iowa Kansas Missouri - Entire State Montana Nebraska Nevada North Dakota Northern Mariana Islands Oregon South Dakota Utah Washington Wyoming

Article Information

General Information

Article ID
A52457

Original ICD-9 Article ID
A19885
A47227
A19806
A19800

Article Title
Ankle-Foot/Knee-Ankle-Foot Orthoses - Policy Article

Article Type
Article

AMA CPT / ADA CDT / AHA NUBC Copyright Statement

Original Effective Date
10/01/2015

Revision Effective Date
10/01/2020

Revision Ending Date
N/A

Retirement Date
N/A

Article Guidance

Article Text:

NON-MEDICAL NECESSITY COVERAGE AND PAYMENT RULES

For any item to be covered by Medicare, it must 1) be eligible for a defined Medicare benefit category, 2) be reasonable and necessary for the diagnosis or treatment of illness or injury or to improve the functioning of a malformed body member, and 3) meet all other applicable Medicare statutory and regulatory requirements. Information provided in this policy article relates to determinations other than those based on Social Security Act §1862(a)(1)(A) provisions (i.e. "reasonable and necessary").

For a beneficiary's equipment to be eligible for reimbursement the reasonable and necessary (R&N) requirements set out in the related Local Coverage Determination must be met. In addition, there are specific statutory payment policy requirements, discussed below, that also must be met.

Ankle-foot orthoses (AFO) and knee-ankle foot orthoses (KAFO) are covered under the Medicare Braces Benefit

(Social Security Act §1861(s)(9)). For coverage under this benefit, the orthosis must be a rigid or semi-rigid device, which is used for the purpose of supporting a weak or deformed body member or restricting or eliminating motion in a diseased or injured part of the body. Items that are not sufficiently rigid to be capable of providing the necessary immobilization or support to the body part for which it is designed do not meet the statutory definition of the Braces Benefit. Items that do not meet the definition of a brace are statutorily noncovered, no benefit.

Both "off-the-shelf" (OTS) and custom-fit items are considered prefabricated braces for Medicare coding purposes. 42 CFR §414.402 establishes that correct coding of AFO and KAFO items is dependent upon whether there is a need for "minimal self-adjustment" during the final fitting at the time of delivery. (See definitions below in Coding Guidelines). If a custom fit code is billed when minimal self-adjustment was provided at final delivery, or if an OTS code is billed when more than minimal self-adjustments were made at final delivery, the claims will be denied as incorrect coding with a statutory denial.

A static/dynamic Ankle-Foot Orthosis (AFO) (L4396, L4397) and replacement interface (L4392) are denied as noncovered (no Medicare benefit) when they are used solely for the prevention or treatment of a heel pressure ulcer because for these indications they are not used to support a weak or deformed body member or to restrict or eliminate motion in a diseased or injured part of the body (i.e., it does not meet the definition of a brace).

A foot drop splint/recumbent positioning device (L4398) and replacement interface (L4394) are denied as noncovered (no Medicare benefit) when they are used solely for the prevention or treatment of a pressure ulcer because for these indications they are not used to support a weak or deformed body member or to restrict or eliminate motion in a diseased or injured part of the body (i.e., it does not meet the definition of a brace).

Elastic or other fabric support garments (A4467 (BELT, STRAP, SLEEVE, GARMENT, OR COVERING, ANYTYPE)) with or without stays or panels do not meet the statutory definition of a brace because they are not rigid or semi-rigid devices. Code A4467 is denied as noncovered (no Medicare benefit). Refer to the coding guideline below for additional information.

A foot pressure off-loading/supportive device (A9283) is denied as noncovered (no Medicare benefit), because it does not support a weak or deformed body member or restrict or eliminate motion in a diseased or injured part of the body.

An inversion/eversion correction device (A9285) is denied as noncovered (no Medicare benefit), because it does not act as a brace; that is, it does not support a weak or deformed body member or restrict or eliminate motion in a diseased or injured part of the body.

Socks (L2840, L2850) used in conjunction with orthoses are denied as noncovered (no Medicare benefit).

Refer to the Orthopedic Footwear policy for information on coverage of shoes and related items which are an integral part of a brace.

There is no separate payment if CAD-CAM technology is used to fabricate an orthosis. Reimbursement is included in the allowance of the codes for custom fabricated orthoses.

Evaluation of the beneficiary, measurement and/or casting, and fitting/adjustments of the orthosis are included in the allowance for the orthosis. There is no separate payment for these services.

Payment for ankle-foot orthoses or knee-ankle foot orthoses are included in the payment to a hospital or skilled nursing facility (SNF) if:

1. The orthosis is provided to a beneficiary prior to an inpatient hospital admission or Part A covered SNF stay;

and,
2. The medical necessity for the orthosis begins during the hospital or SNF stay (e.g., after ankle, foot, or knee surgery).

A claim should not be submitted to the DME MAC in this situation.

Payment for ankle-foot orthoses or knee-ankle foot orthoses are also included in the payment to a hospital or a Part A covered SNF stay if:

1. The orthosis is provided to a beneficiary during an inpatient hospital or Part A covered SNF stay prior to the day of discharge; and,
2. The beneficiary uses the item for medically necessary inpatient treatment or rehabilitation.

A claim must not be submitted to the DME MAC in this situation.

Payment for ankle-foot orthoses or knee-ankle foot orthoses delivered to a beneficiary in a hospital or a Part A covered SNF stay is eligible for coverage by the DME MAC if:

1. The orthosis is medically necessary for a beneficiary after discharge from a hospital or Part A covered SNF stay; and,
2. The orthosis is provided to the beneficiary within two days prior to discharge to home; and,
3. The orthosis is not needed for inpatient treatment or rehabilitation, but is left in the room for the beneficiary to take home.

REQUIREMENTS FOR SPECIFIC DMEPOS ITEMS PURSUANT TO Final Rule 1713 (84 Fed. Reg Vol 217)

Final Rule 1713 (84 Fed. Reg Vol 217) requires a face-to-face encounter and a Written Order Prior to Delivery (WOPD) for specified HCPCS codes. CMS and the DME MACs provide a list of the specified codes, which is periodically updated. The link will be located here once it is available.

Claims for the specified items subject to Final Rule 1713 (84 Fed. Reg Vol 217) that do not meet the face-to-face encounter and WOPD requirements specified in the LCD-related Standard Documentation Requirements Article (A55426) will be denied as not reasonable and necessary.

If a supplier delivers an item prior to receipt of a WOPD, it will be denied as not reasonable and necessary. If the WOPD is not obtained prior to delivery, payment will not be made for that item even if a WOPD is subsequently obtained by the supplier. If a similar item is subsequently provided by an unrelated supplier who has obtained a WOPD, it will be eligible for coverage.

POLICY SPECIFIC DOCUMENTATION REQUIREMENTS

In addition to policy specific documentation requirements, there are general documentation requirements that are applicable to all DMEPOS policies. These general requirements are located in the DOCUMENTATION REQUIREMENTS section of the LCD.

Refer to the LCD-related Standard Documentation Requirements article, located at the bottom of this Policy Article

under the Related Local Coverage Documents section for additional information regarding GENERAL DOCUMENTATION REQUIREMENTS and the POLICY SPECIFIC DOCUMENTATION REQUIREMENTS discussed below.

General Requirements

The supplier must include on the claim line the diagnosis code(s) for HCPCS codes L4396, L4397, L4392 and L4631.

For a custom-fabricated orthosis, there must be documentation in the supplier's records to support the medical necessity of that type device rather than a prefabricated orthosis. This information must be available upon request.

When providing these items suppliers must:

- Provide the product that is specified by the treating practitioner
- Be sure that the treating practitioner's medical record justifies the need for the type of product (i.e., Prefabricated versus Custom Fabricated)
- Only bill for the HCPCS code that accurately reflects both the type of orthosis and the appropriate level of fitting
- Have detailed documentation in the supplier's record that justifies the code selected

For prefabricated orthoses (L1902, L1906, L1910, L1930, L1932, L1951, L1971, L2035, L2112, L2114, L2116, L2132, L2134, L2136, L4350, L4360, L4361, L4370, L4386, L4387, L4396, L4397, L4398), there is no physical difference between orthoses coded as custom fitted versus those coded as off-the-shelf. The differentiating factor for proper coding (see definitions in Coding Guidelines below) is the need for “minimal self-adjustment” at the time of fitting by the beneficiary, caretaker for the beneficiary, or supplier. This minimal self-adjustment does not require the services of a certified orthotist or an individual who has specialized training. Items requiring minimal self-adjustment are coded as off-the-shelf orthoses. For example, adjustment of straps and closures, bending or trimming for final fit or comfort (not all-inclusive) fall into this category.

Fabrication of an orthosis using CAD/CAM or similar technology without the creation of a positive model with minimal self-adjustment at delivery is considered as OTS.

Items requiring more than minimal self-adjustment by a qualified practitioner (as defined in the Coding Guidelines below) are coded as custom fitted (L1910, L1930, L1932, L1951, L1971, L2035, L2112, L2114, L2116, L2132, L2134, L2136, L4360, L4386, L4396). Documentation must be sufficiently detailed to include, but is not limited to, a detailed description of the modifications necessary at the time of fitting the orthosis to the beneficiary. This information must be available upon request.

For custom fabricated orthoses (L1904, L1907, L1920, L1940, L1945, L1950, L1960, L1970, L1980, L1990, L2000, L2005, L2006, L2010, L2020, L2030, L2034, L2036, L2037, L2038, L2106, L2108, L2126, L2128, L4631), there must be detailed documentation in the treating practitioner's records to support the medical necessity of custom fabricated rather than a prefabricated orthosis as described in the Coverage Indications, Limitations and/or Medical Necessity section of the related LCD. This information will be corroborated by the functional evaluation in the orthotist or prosthetist’s records. This information must be available upon request.

MODIFIERS

KX, GA, and GZ MODIFIERS:

Suppliers must add a KX modifier to the AFO/KAFO base and addition codes only if all of the coverage criteria in the

"Coverage Indications, Limitations and or Medical Necessity" section in the related LCD have been met and evidence of such is retained in the supplier's files and available to the DME MAC upon request.

If all of the criteria in the Coverage Indications, Limitations and/or Medical Necessity section of the related LCD have not been met, the GA or GZ modifier must be added to the code. When there is an expectation of a medical necessity denial, suppliers must enter the GA modifier on the claim line if they have obtained a properly executed Advance Beneficiary Notice (ABN) or the GZ modifier if they have not obtained a valid ABN.

Claims lines billed with codes without a KX, GA or GZ modifier will be rejected as missing information.

MISCELLANEOUS

If the item is custom fabricated and does not have a specific HCPCS code, a complete and clear description of the item, including what makes this item unique, and a breakdown of charges (material and labor used in fabrication) should be entered in the narrative field of an electronic claim or on Item 19 of a paper claim. (Refer to the LCD-related Standard Documentation Requirements article (A55426) for more information regarding billing of items with HCPCS codes that include miscellaneous, NOC, unlisted, or non-specified in their narrative descriptions.)

A claim for code L4205 must include an explanation of what is being repaired. A claim for code L4210 must include a description of each item that is billed. This information should be entered in the narrative field of an electronic claim.

All codes for orthoses or repairs of orthoses billed with the same date of service must be submitted on the same claim.

Refer to the Orthopedic Footwear policy for information on documentation requirements for shoes and related items which are an integral part of a brace.

CODING GUIDELINES

Off-the-shelf (OTS) orthotics are:

- Items that are prefabricated.
- They may or may not be supplied as a kit that requires some assembly. Assembly of the item and/or installation of add-on components and/or the use of some basic materials in preparation of the item does not change classification from OTS to custom fitted.
- OTS items require minimal self-adjustment for fitting at the time of delivery for appropriate use and do not require expertise in trimming, bending, and molding, assembling, or customizing to fit an individual.
- This fitting does not require expertise of a certified orthotist or an individual who has specialized training in the provision of orthoses to fit the item to the individual beneficiary.

The term "minimal self-adjustment" is defined at 42 CFR §414.402 as an adjustment the beneficiary, caregiver for the beneficiary, or supplier of the device can perform and that does not require the services of a certified orthotist (that is, an individual who is certified by the American Board for Certification in Orthotics and Prosthetics, Inc., or by the Board for Orthotist/Prosthetist Certification) or an individual who has specialized training. For example, adjustment of straps and closures, bending or trimming for final fit or comfort (not all-inclusive) fall into this category. See "more than minimal self-adjustment" definition below for additional information.

Use of CAD/CAM or similar technology to create an orthosis without a positive model of the patient may be considered as OTS if the final fitting upon delivery to the patient requires minimal self-adjustment not requiring expertise as described in this section.

Custom fitted orthotics are:

- Devices that are prefabricated.
- They may or may not be supplied as a kit that requires some assembly. Assembly of the item and/or installation of add-on components and/or the use of some basic materials in preparation of the item does not change classification from OTS to custom fitted.
- Classification as custom fitted requires more than minimal self-adjustment for fitting at the time of delivery in order to provide an individualized fit, i.e., the item must be trimmed, bent, molded (with or without heat), or otherwise modified resulting in alterations beyond minimal self-adjustment.
- This fitting at delivery does require expertise of a certified orthotist or an individual who has specialized training in the provision of orthosis to fit the item to the individual beneficiary.

In contrast to "minimal self-adjustment," "more than minimal self-adjustment" is defined as changes made to achieve an individualized fit during the final fitting at the time of delivery of the item that requires the expertise of a certified orthotist or an individual who has equivalent specialized training in the provision of orthotics in compliance with all applicable Federal and State licensure and regulatory requirements. A certified orthotist is defined as an individual who is certified by the American Board for Certification in Orthotics and Prosthetics, Inc., or by the Board for Orthotist/Prosthetist Certification.

In most cases for prefabricated orthoses, the correct coding of the orthosis is dictated by actions that take place at the time of fitting to the beneficiary, either custom-fit (requiring expertise) or OTS (requiring minimal self-adjustment). However, for certain types of orthoses, the HCPCS code narrative that best describes the product does not make a distinction between prefabricated orthoses that are provided as custom-fit or OTS. These code narratives are correct and must be used for Medicare billing, without regard to how the product is provided to the beneficiary at the final delivery.

There are products that may be either fit by the beneficiary or require custom fitting at the time of final delivery. There are parallel sets of HCPCS codes (L4360, L4361, L4386, L4387, L4396 and L4397) that describe identical types of items. The codes are only differentiated based upon the nature of the final fitting performed at the time of delivery. The alternative HCPCS code types are:

- HCPCS codes which describe "PREFABRICATED, OFF-THE-SHELF" must be used when minimal self-adjustment is the extent of the fitting performed at delivery.
- HCPCS codes which describe "PREFABRICATED ITEM THAT HAS BEEN TRIMMED, BENT, MOLDED, ASSEMBLED, OR OTHERWISE CUSTOMIZED TO FIT A SPECIFIC PATIENT BY AN INDIVIDUAL WITH EXPERTISE" must be used when more than minimal self-adjustment is necessary at delivery.

Use of CAD/CAM or similar technology to create an orthosis without a positive model of the patient may be considered as custom fitted if the final fitting at the time of delivery to the patient requires more than minimal self-adjustment requiring expertise as described in this section.

Kits are:

- A collection of components, materials, and parts that require further assembly before delivery of the final product.
- The elements of a kit may be packaged and complete from a single source or may be an assemblage of separate components from multiple sources by the supplier.

Elastic and Similar Stretchable Materials

For items where the HCPCS code specifies “elastic” or other similar terminology for stretchable material, use the code that is most applicable to the item. A NOC (Not Otherwise Classified) or miscellaneous HCPCS code must not be used instead of the specific code. Refer to the long code narrative and any relevant coding guideline for the criteria applicable for each HCPCS code.

For items where the HCPCS code does not specify elastic or other similar terminology for stretchable material, the following guidelines apply:

- Items that are primarily constructed of elastic or other stretchable materials (e.g. support items made of material such as neoprene or spandex (elastane, Lycra®) (not all-inclusive)) must be coded as A4467 (BELT, STRAP, SLEEVE, GARMENT, OR COVERING, ANY TYPE).
- Items that are primarily constructed of elastic or other stretchable materials (e.g. support items made of material such as neoprene or spandex (elastane, Lycra®]) (not all-inclusive)) that contain stays and/or panels must be coded as A4467 (BELT, STRAP, SLEEVE, GARMENT, OR COVERING, ANY TYPE).
- Items that are primarily constructed of inelastic material (e.g., canvas, cotton or nylon (not all-inclusive)) that are incapable of providing the necessary immobilization or support to the body part for which it is designed must be coded using A4467 (BELT, STRAP, SLEEVE, GARMENT, OR COVERING, ANY TYPE).
- Items that are primarily of constructed inelastic material (e.g., canvas, cotton or nylon (not all-inclusive)) that are incapable of providing the necessary immobilization or support to the body part for which it is designed and that have stays and/or panels capable of providing the required immobilization or support to the body part for which it is designed, must be coded using A4467 (BELT, STRAP, SLEEVE, GARMENT, OR COVERING, ANY TYPE).
- Items that are primarily constructed of inelastic material (e.g., canvas, cotton or nylon (not all-inclusive)) capable of providing the necessary immobilization or support to the body part for which it is designed must be coded using the applicable specific HCPCS code for the type of product. A NOC (Not Otherwise Classified) or miscellaneous HCPCS code must not be used instead of the specific code. Refer to the long code narrative and any relevant coding guideline for the criteria applicable for each HCPCS code.
- Items that are primarily of constructed inelastic material (e.g., canvas, cotton or nylon (not all-inclusive)) capable of providing the necessary immobilization or support to the body part for which it is designed and that have stays and/or panels capable of providing the required immobilization or support to the body part for which it is designed, must be coded using the applicable specific HCPCS code for the type of product. A NOC (Not Otherwise Classified) or miscellaneous HCPCS code must not be used instead of the specific code. Refer to the long code narrative and relevant coding guideline for the criteria applicable for each HCPCS code.
- Items that are not capable of providing the necessary immobilization or support to the body part for which it is designed (regardless of materials) must be coded using A9270 (NONCOVERED ITEM OR SERVICE).

Ankle-foot orthoses described by codes L1900, L1910, L1920, L1930, L1932, L1940, L1945, L1950, L1951, L1960, L1970, L1971, L1980, L1990, extend well above the ankle (usually to near the top of the calf) and are fastened around the lower leg above the ankle. These features distinguish them from foot orthotics which are shoe inserts that do not extend above the ankle and ankle gauntlets described by codes L1902, L1904, L1906, L1907.

L1900 (ANKLE FOOT ORTHOSIS, SPRING WIRE, DORSIFLEXION ASSIST CALF BAND, CUSTOM FABRICATED) describes a custom fabricated Ankle Foot Orthosis (AFO) designed to control inversion, eversion, dorsiflexion, and plantarflexion motions of the ankle foot complex. Primary construction is of two springwire uprights, joined to rigid calf band/cuff, and component for attaching to the sole of an orthopedic shoe. Springwire will be contoured into a coiled spring below the ankle and above shoe sole. Height of rigid calf band/cuff terminates well above the ankle (usually to near the top of the calf) and is fastened around the lower leg above the ankle. The coiled springwire applies a spring force to resist plantarflexion motion and provide dorsiflexion assist. Included in the code are closure components, and for attaching spring wire to footwear. This AFO is custom fabricated per the DMEPOS quality

standards, Appendix C.

L1902 (ANKLE ORTHOSIS, ANKLE GAUNTLET OR SIMILAR, WITH OR WITHOUT JOINTS, PREFABRICATED, OFF-THE-SHELF) describes a prefabricated Ankle Orthosis designed to provide compression and resist motion of the ankle foot complex. Primary construction is a sleeve type device (gauntlet) with or without joints. The gauntlet encloses the foot and ankle from the longitudinal arch to at least just above the malleoli. Gauntlet closure may be straps, hook and loop, laces or equal. May or may not include joints or hinges for additional support. There are no additional HCPCS codes for this type of prefabricated ankle orthosis.

L1904 (ANKLE ORTHOSIS, ANKLE GAUNTLET OR SIMILAR, WITH OR WITHOUT JOINTS, CUSTOM FABRICATED) describes a custom fabricated Ankle Orthosis designed to provide compression and resist motion of the ankle foot complex. Primary construction is a sleeve type device (gauntlet) with or without joints. The gauntlet encloses the foot and ankle from the longitudinal arch to at least just above the malleoli. Gauntlet closure may be straps, hook and loop, laces or equal. May or may not include joints or hinges for additional support. Ankle Orthosis is custom fabricated per the DMEPOS quality standards, Appendix C.

L1906 (ANKLE FOOT ORTHOSIS, MULTILIGAMENTOUS, ANKLE SUPPORT, PREFABRICATED, OFF-THE-SHELF) describes a prefabricated Ankle Foot Orthosis (AFO) which provides multi-directional support to the ankle ligaments while allowing free dorsiflexion and plantarflexion motion. Primary construction contains a rigid footplate with integral ankle joints and uprights extending above and below the ankle joints. Proximal uprights provide surface area contact for stabilization of the footplate and ankle joints. Additional support is from wraparound straps. Included in the code are closures from lacing, webbing, hook and loop, or equal, and additional support strapping. There are no additional HCPCS codes for this type of prefabricated ankle orthosis. Effective for claims with dates of service on or after April 1, 2012, the only products which may be billed to Medicare using code L1906 are those for which a written coding verification has been made by the Pricing, Data Analysis, and Coding (PDAC) contractor and that are listed in the Product Classification List.

L1907 (ANKLE ORTHOSIS, SUPRAMALLEOLAR WITH STRAPS, WITH OR WITHOUT INTERFACE/PADS, CUSTOM FABRICATED) describes a custom fabricated Ankle Orthosis (AO) designed to control motion of the ankle and mid-foot. Primary construction is of molded plastic designed to control inversion and eversion, and horizontal rotation motions while allowing dorsiflexion and plantarflexion motion of the ankle joint. Included in the code are closure components, soft interface and padding. Trim lines extend from tips of toes to just above the malleoli. AO is custom fabricated per the DMEPOS quality standards, Appendix C.

L1910 (ANKLE FOOT ORTHOSIS, POSTERIOR, SINGLE BAR, CLASP ATTACHMENT TO SHOE COUNTER, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT) describes a prefabricated Ankle Foot Orthosis (AFO) designed to control dorsiflexion and plantarflexion motions of the ankle foot complex. Primary construction is a single flexible upright utilizing a removable clasp-type component attached to the heel portion of a shoe and a rigid calf band/cuff. Height of the rigid calf band/cuff terminates well above the ankle (usually to near the top of the calf) and is fastened around the lower leg above the ankle. Included in the code are closure and clasp components.

L1920 (ANKLE FOOT ORTHOSIS, SINGLE UPRIGHT WITH STATIC OR ADJUSTABLE STOP (PHELPS OR PERLSTEIN TYPE), CUSTOM FABRICATED) describes a custom fabricated Ankle Foot Orthosis designed to control only the plantarflexion motion of the ankle foot complex. Primary construction is a single metal upright which has a pivoting attachment into a sole component attached to an orthopedic shoe. The single rigid upright is joined to a rigid calf band/cuff which terminates well above the ankle (usually to near the top of the calf) and is fastened around the lower leg above the ankle. The pivoting attachment has an integrated component(s) to stop plantarflexion ankle motion while always allowing free dorsiflexion motion. Included in the code are components for closures and attaching the footwear. This AFO is custom fabricated per the DMEPOS quality standards, Appendix C.

L1930 (ANKLE FOOT ORTHOSIS, PLASTIC OR OTHER MATERIAL, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT) describes a prefabricated Ankle Foot Orthosis designed to control inversion, eversion, dorsiflexion, and plantarflexion motions of the ankle foot complex. Primary construction is a rigid calf cuff and rigid foot plate section joined by a flexible posterior strut. Foot plate must extend to the metatarsal heads and may extend as far as the toe tips. Calf cuff height terminates well above the ankle (usually to near the top of the calf) and are fastened around the lower leg above the ankle and footplate may extend to toe tip. This AFO is constructed from plastic or other materials. Included in the code are closure components.

L1932 (AFO, RIGID ANTERIOR TIBIAL SECTION, TOTAL CARBON FIBER OR EQUAL MATERIAL, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT) describes a prefabricated Ankle Foot Orthosis designed to control the dorsiflexion and plantarflexion, and inversion and eversion, motions of the ankle foot complex. Primary construction includes full length foot plate, rigid front shin shell that extends from lower shin region to near tibial turbercle. Rigid strut connects foot plate to shin shell. This AFO is constructed of carbon fiber or equal. Included in the code are closure components and soft interface.

L1940 (ANKLE FOOT ORTHOSIS, PLASTIC OR OTHER MATERIAL, CUSTOM FABRICATED) describes a custom fabricated Ankle Foot Orthosis designed to control inversion, eversion, dorsiflexion, and plantarflexion motions of the ankle foot complex. Primary construction includes rigid foot plate, strut or equal component, which joins footplate to a calf cuff. Calf cuff height terminates well above the ankle (usually to near the top of the calf) and are fastened around the lower leg above the ankle. This AFO can be constructed from flexible and strong thermosetting materials, thermoplastics, or composite type materials. Included in the code are closure components. AFO is custom fabricated per the DMEPOS quality standards, Appendix C.

L1945 (ANKLE-FOOT ORTHOSIS (AFO), PLASTIC, RIGID ANTERIOR TIBIAL SECTION (FLOOR REACTION), CUSTOM FABRICATED) describes a custom fabricated Ankle Foot Orthosis designed to control inversion, eversion, dorsiflexion, plantarflexion, and horizontal rotation motions of the ankle foot complex. Primary construction is a plastic full-length rigid foot plate, rigid shell with Anterior calf cuff typically extending from mid-shin region to tibial tubercle. Rigid strut connects foot plate to shin shell. Shell cutouts are created to aid donning of the AFO. This AFO is constructed of plastic materials. Included in the code are closure components. AFO is custom fabricated per the DMEPOS quality standards, Appendix C.

L1950 (ANKLE FOOT ORTHOSIS, SPIRAL, (INSTITUTE OF REHABILITATIVE MEDICINE TYPE), PLASTIC, CUSTOM FABRICATED) describes a custom fabricated Ankle Foot Orthosis designed to control inversion, eversion, dorsiflexion, plantarflexion, and horizontal rotation motions of the ankle foot complex. Primary construction includes, at minimum, a 90-degree spiral-shaped strut joining the rigid footplate to the rigid calf cuff. Foot plate must extend to the metatarsal heads and may extend as far as the toe tips. Calf cuff height terminates well above the ankle (usually to near the top of the calf) and are fastened around the lower leg above the ankle. This AFO can be constructed from flexible and strong thermosetting materials, thermoplastics, or composite type materials. Included in the code are closure components. AFO is custom fabricated per the DMEPOS quality standards, Appendix C.

L1951 (ANKLE FOOT ORTHOSIS, SPIRAL, (INSTITUTE OF REHABILITATIVE MEDICINE TYPE), plastic or other material, prefabricated, includes fitting and adjustment) describes a prefabricated Ankle Foot Orthosis designed to control dorsiflexion and plantarflexion motions of the ankle foot complex. Primary construction includes, at minimum, a 90-degree spiral-shaped strut linking the footplate to the rigid calf cuff. Foot plate must extend to the metatarsal heads and may extend as far as the toe tips. Calf cuff height terminates well above the ankle (usually to near the top of the calf) and are fastened around the lower leg above the ankle. The AFO can be constructed from flexible and strong thermosetting materials, thermoplastics, or composite type materials. Included in the code are closure components.

L1960 describes an Ankle Foot Orthosis (AFO) which provides ankle control for beneficiaries with musculoskeletal or neuromuscular dysfunction. The AFO is designed to provide rigid immobilization of the ankle-foot complex in the sagittal, coronal, and transverse planes. The custom fabricated solid ankle AFO can be constructed from thermosetting materials, thermoplastics, or composite type materials.

L1970 (ANKLE FOOT ORTHOSIS, PLASTIC WITH ANKLE JOINT, CUSTOM FABRICATED**)** describes a custom fabricated Ankle Foot Orthosis designed to control inversion, eversion, dorsiflexion, plantarflexion, and horizontal rotation motions of the ankle foot complex. Primary construction is a rigid shell-like or equal structure containing a hinge or joint mechanism. Structure contacts calf, lower leg, and foot to provide control. Foot plate must extend to the metatarsal heads and may extend as far as the toe tips. Calf cuff height terminates well above the ankle (usually to near the top of the calf) and are fastened around the lower leg above the ankle. This AFO can be constructed from thermosetting materials, thermoplastics, or composite type materials. Included in the code are closure components. AFO is custom fabricated per the DMEPOS quality standards, Appendix C.

L1971 (ANKLE FOOT ORTHOSIS, PLASTIC OR OTHER MATERIAL WITH ANKLE JOINT, PREFABRICATED, INCLUDES FITTING AND ADJUSTMENT) describes a prefabricated Ankle Foot Orthosis designed to control inversion, eversion, dorsiflexion, plantarflexion, and horizontal rotation motions of the ankle foot complex. Primary construction is a rigid shell-like or equal structure containing a hinge or joint mechanism. Structure contacts calf, lower leg, and foot to provide control. Foot plate must extend to the metatarsal heads and may extend as far as the toe tips. Calf cuff height terminates well above the ankle.

L1980 (ANKLE FOOT ORTHOSIS, SINGLE UPRIGHT FREE PLANTAR DORSIFLEXION, SOLID STIRRUP, CALF BAND/CUFF (SINGLE BAR 'BK' ORTHOSIS), CUSTOM FABRICATED) describes a custom fabricated Ankle Foot Orthosis designed to control inversion and eversion motions of the ankle foot complex. Primary construction is a single metal upright (medial or lateral) joined to a rigid calf band, free motion ankle joint, and stirrup component attached to an orthopedic shoe. Height of calf cuff/band terminates well above the ankle (usually to near the top of the calf) and is fastened around the lower leg above the ankle. Included in the code are components for closures and attaching the footwear. This AFO is custom fabricated per the DMEPOS quality standards, Appendix C.

L1990 (ANKLE FOOT ORTHOSIS, DOUBLE UPRIGHT FREE PLANTAR DORSIFLEXION, SOLID STIRRUP, CALF BAND/CUFF (DOUBLE BAR 'BK' ORTHOSIS), CUSTOM FABRICATED) describes a custom fabricated Ankle Foot Orthosis designed to control inversion and eversion motions of the ankle foot complex. Primary construction are two metal uprights (medial and lateral) joined to a rigid calf band, free motion ankle joints, and stirrup component attached to an orthopedic shoe. Height of calf cuff/band terminates well above the ankle (usually to near the top of the calf) and is fastened around the lower leg above the ankle. Included in the code are components for closures and attaching the footwear. This AFO is custom fabricated per the DMEPOS quality standards, Appendix C.

L2006 describes a custom fabricated, single or double upright Knee-Ankle Foot Orthosis (KAFO) with an adjustable microprocessor control feature which provides resistance to stance and swing phase knee joint motion. The custom fabricated KAFO can be constructed from thermosetting materials, thermoplastics, or composite type materials. There are no additional add-on codes for this KAFO.

Effective for claims with dates of service on or after January 1, 2020, the only products which may be billed to Medicare using code L2006 are those for which a written coding verification has been made by the Pricing, Data Analysis, and Coding (PDAC) contractor and that are listed in the Product Classification List.

L2340 is a pre-tibial shell, custom fabricated, that provides a rigid overlapping interlocking anterior tibial control between the tibial tuberosity to a point no greater than 3 inches proximal to the medial malleolus. The pre-tibial shell can be constructed from thermosetting materials, thermoplastics, or composite type materials.

Code L2755 describes an addition to a lower extremity orthosis composed of high strength and/or lightweight material such as Kevlar®, carbon fiber or other laminated or impregnated composite material.

A nonambulatory ankle-foot orthosis may be either an ankle contracture splint, night splint or a foot drop splint.

A static or dynamic positioning ankle-foot orthosis (L4396, L4397) is a prefabricated ankle-foot orthosis which has all of the following characteristics:

1. Designed to accommodate either plantar fasciitis or an ankle with a plantar flexion contracture up to 45°; and,
2. Applies a dorsiflexion force to the ankle; and,
3. Used by a beneficiary who is minimally ambulatory, or nonambulatory; and,
4. Has a soft interface.

A foot drop splint/recumbent positioning device (L4398) is a prefabricated ankle-foot orthosis which has all of the following characteristics:

1. Designed to maintain the foot at a fixed position of 0° (i.e., perpendicular to the lower leg); and,
2. Not designed to accommodate an ankle with a plantar flexion contracture; and,
3. Used by a beneficiary who is nonambulatory; and,
4. Has a soft interface.

Code L4631 describes a Charcot's restraint orthotic walker (CROW) orthosis. Code L4631 is a custom fabricated ankle-foot orthosis which has all of the following characteristics:

1. Designed to maintain the foot at a fixed position of 0° (i.e., perpendicular to the lower leg); and,
2. Allows for varus or valgus deformity correction; and,
3. Contains a rocker bottom sole with a custom arch support; and,
4. Incorporates a rigid anterior tibial shell; and,
5. Used by a beneficiary who is ambulatory; and,
6. Has a soft interface
7. Totally encapsulated.

Code L4631 includes all additions including straps and closures. No additional codes may be billed with code L4631.

Codes L1900, L1904, L1907, L1920, L1940, L1945, L1950, L1960, L1970, L1980, L1990, L2000, L2005, L2006, L2010, L2020, L2030, L2034, L2036, L2037, L2038, L2106, L2108, L2126, L2128 and L4631 describe custom-fabricated orthoses. These codes must not be used for prefabricated orthoses.

Codes L1902, L1906, L1910, L1930, L1932, L1951, L1971, L2035, L2112, L2114, L2116, L2132, L2134, L2136, L4350, L4360, L4361, L4370, L4386, L4387, L4392, L4394, L4396, L4397, and L4398 describe prefabricated orthoses. These codes must not be used for custom-fabricated orthoses.

Codes L1900, L1902, L1904, L1906, L1907, L1910, L1920, L1930, L1932, L1940, L1945, L1950, L1951, L1960, L1970, L1971, L1980, L1990, L2106, L2108, L2112, L2114, L2116, L4350, L4360, L4361, L4386, L4387 and L4631 are used for an ankle-foot orthosis which is worn when a beneficiary is ambulatory.

Codes L4396 and L4397 are used for an ankle-foot orthosis which is worn when a beneficiary is nonambulatory, or minimally ambulatory.

Code L4398 is used for an ankle-foot orthosis which is worn when a beneficiary is nonambulatory.

Some replacement items have unique Healthcare Common Procedure Coding System (HCPCS) codes. Replacement components that do not have a unique HCPCS code must be billed with a "not otherwise specified" code - L2999. Items that have unique codes must not be billed using a NOC code.

HCPCS codes L4050 and L4055 do not describe replacement soft interfaces used with contracture orthoses.

Foot orthotics are shoe inserts that do not extend above the ankle. The correct codes for foot orthotics provided for beneficiaries without diabetes are L3000, L3001, L3002, L3003, L3010, L3020, L3030, L3031, L3040, L3050, L3060, L3070, L3080, L3090 (Refer to the Orthopedic Footwear policy for more information). Multiple density foot orthotics used in the management of diabetic foot problems are coded A5512, A5513, and A5514 (code A5514 effective for DOS on or after 01/01/2019) (Refer to the Therapeutic Shoes for Persons with Diabetes policy for more information).

All claims for devices that contain a concentric adjustable torsion style mechanism in the knee joint for any condition other than an assistive function to joint extension motion must be coded as Durable Medical Equipment using code E1810 (DYNAMIC ADJUSTABLE KNEE EXTENSION / FLEXION DEVICE, INCLUDES SOFT INTERFACE MATERIAL). If a concentric adjustable torsion style mechanism in the knee joint is used solely to provide an assistive function for joint extension, it must be coded as L2999 (See Coverage Indications, Limitations and/or Medical Necessity section of the related LCD).

All claims for devices that contain a concentric adjustable torsion style mechanism in the ankle joint for any condition other than an assistive function to joint plantar- or dorsiflexion motion must be coded as Durable Medical Equipment using code E1815 (DYNAMIC ADJUSTABLE ANKLE EXTENSION/FLEXION DEVICE, INCLUDES SOFT INTERFACE MATERIAL). If a concentric adjustable torsion style mechanism in the ankle joint is used solely to provide an assistive function for joint plantar or dorsiflexion, it must be coded as L2999 (See Coverage Indications, Limitations and/or Medical Necessity section of the related LCD).

Claims for devices that contain a concentric adjustable torsion style mechanism in the knee or ankle joint and that are being used to treat any condition other than an assistive function to joint extension motion are not covered under the Braces benefit and will be denied as incorrect coding when billed using code L2999 (See Coverage Indications, Limitations and/or Medical Necessity section of the related LCD).

Code A9283 (FOOT PRESSURE OFF LOADING/SUPPORTIVE DEVICE, ANY TYPE, EACH) is used for an item that is designed primarily to reduce pressure on the sole or heel of the foot. It may be a shoe-like item, an item that is used inside a shoe and may or may not extend outside the shoe, or an item that is attached to a shoe. It may be prefabricated or custom fabricated. Code A9283 does not include items that meet the definition of a therapeutic shoe for diabetes (A5500, A5501).

Prefabricated walking boots are coded using codes L4360, L4361, L4386 or L4387. These codes describe complete products. Claims for add-on codes used with walking boots coded L4360, L4361, L4386 or L4387 will be denied as unbundling.

Certain products may have both covered and non-covered uses, as defined by the Braces benefit category, and must be coded based on the beneficiary’s condition. For example, when used as a brace for the treatment of an orthopedic condition, walking boots are coded L4360, L4361, L4386, L4387 and L4631. However, walking boots must be coded A9283 when used solely for the prevention or treatment of a lower extremity ulcer or pressure reduction.

When using code A9283, there is no separate billing using addition codes. Replacement liners for devices billed with A9283 must be billed with code A9270 (noncovered item or service).

Code A9285 (INVERSION/EVERSION CORRECTION DEVICE) is designed to provide off-loading pressure to the knee for the treatment of knee osteoarthritis. The device is applied at the foot and extends across the ankle to apply pressure to the side of the leg below the knee. It does not provide any support at the ankle.

The right (RT) and left (LT) modifiers must be used with orthosis base codes, additions, and replacement parts. Effective for claims with dates of service (DOS) on or after 3/1/2019, when the same code for bilateral items (left and right) is billed on the same date of service, bill each item on two separate claim lines using the RT and LT modifiers and 1 unit of service (UOS) on each claim line. Do not use the RTLT modifier on the same claim line and billed with 2 UOS. Claims billed without modifiers RT and/or LT, or with RTLT on the same claim line and 2 UOS, will be rejected as incorrect coding.

Code L4205 (Repair of orthotic device, labor component, per 15 minutes) may only be billed for time involved with the actual repair of an orthosis or for medically necessary adjustments made more than 90 days after delivery. Code L4205 must not be used to bill for time involved with other professional services including, but not limited to:

- Evaluating the beneficiary
- Taking measurements, making a cast, making a model, use of CAD/CAM
- Making modifications to a prefabricated item to fit it to the individual beneficiary
- Follow-up visits
- Making adjustments at the time of or within 90 days after delivery

Suppliers must distinguish between repair and replacement of an orthosis. When an orthotic is replaced, there is no separate billing for the above services because reimbursement for these services is included in the allowance for the replacement item.

Repairs to a covered orthosis due to wear or to accidental damage are covered when they are necessary to make the orthosis functional. The reason for the repair must be documented in the supplier's record. If the expense for repairs exceeds the estimated expense of providing another entire orthosis, no payment will be made for the amount in excess.

The allowance for the labor involved in replacing an orthotic component that is coded with a specific L code is included in the allowance for that component. The allowance for the labor involved in replacing an orthotic component that is coded with the miscellaneous code L4210 is separately payable in addition to the allowance for that component.

Addition codes L4002, L4010, L4020, L4030, L4040, L4045, L4050, L4055, L4060, L4070, L4080, L4090, L4100, L4110, L4130, and L4392 are for billing of replacement components and are not payable at initial issue of a base orthosis. When claims for code(s) L4002, L4010, L4020, L4030, L4040, L4045, L4050, L4055, L4060, L4070, L4080, L4090, L4100, L4110, L4130, and L4392 are billed at the time of initial issue of a base orthosis, the addition code(s) will be rejected as incorrect coding.

Suppliers should contact the PDAC contractor for guidance on the correct coding of these items.

Coding Information

CPT/HCPCS Codes

N/A

ICD-10 Codes that Support Medical Necessity

Group 1 Paragraph:

The presence of an ICD-10 code listed in this section is not sufficient by itself to assure coverage. Refer to the LCD section on "**Coverage Indications, Limitations, and/or Medical Necessity**" for other coverage criteria and payment information.

For HCPCS codes L4392, L4396 and L4397:

Group 1 Codes:

ICD-10 CODE	DESCRIPTION
M24.571	Contracture, right ankle
M24.572	Contracture, left ankle
M24.574	Contracture, right foot
M24.575	Contracture, left foot
M72.2	Plantar fascial fibromatosis

Group 2 Paragraph:

For HCPCS code L4631:

Group 2 Codes:

ICD-10 CODE	DESCRIPTION
A52.16	Charcot's arthropathy (tabetic)
E08.610	Diabetes mellitus due to underlying condition with diabetic neuropathic arthropathy
E09.610	Drug or chemical induced diabetes mellitus with diabetic neuropathic arthropathy
E10.610	Type 1 diabetes mellitus with diabetic neuropathic arthropathy
E11.610	Type 2 diabetes mellitus with diabetic neuropathic arthropathy
M14.671	Charcot's joint, right ankle and foot
M14.672	Charcot's joint, left ankle and foot

ICD-10 Codes that DO NOT Support Medical Necessity

Group 1 Paragraph:

For the specific HCPCS codes indicated above, all ICD-10 codes that are not specified in the preceding section. For all other HCPCS codes, diagnoses are not specified.

Group 1 Codes:

N/A

Additional ICD-10 Information

N/A

Bill Type Codes:

Contractors may specify Bill Types to help providers identify those Bill Types typically used to report this service. Absence of a Bill Type does not guarantee that the policy does not apply to that Bill Type.Complete absence of all Bill Types indicates that coverage is not influenced by Bill Type and the policy should be assumed to apply equally to all claims.

N/A

Revenue Codes:

Contractors may specify Revenue Codes to help providers identify those Revenue Codes typically used to report this service. In most instances Revenue Codes are purely advisory. Unless specified in the policy, services reported under other Revenue Codes are equally subject to this coverage determination. Complete absence of all Revenue Codes indicates that coverage is not influenced by Revenue Code and the policy should be assumed to apply equally to all Revenue Codes.

N/A

Revision History Information

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION
10/01/2020	R10	Revision Effective Date: 10/01/2020 MISCELLANEOUS: Clarified: Custom fabricated items that do not have specific HCPCS codes require additional information in claim narrative Added: Statement referring to the LCD-related Standard Documentation Requirements article for more information ICD-10 CODES THAT SUPPORT MEDICAL NECESSITY: Removed: Non-specific ICD-10 codes M24.573 and M24.576 from Group 1 codes Removed: Non-specific ICD-10 code M14.679 from Group 2 codes *09/24/2020: At this time 21st Century Cures Act applies to new and revised LCDs which require comment and notice. This revision is to an article that is not a local coverage determination.*

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION
07/01/2020	R9	Revision Effective Date: 07/01/2020 ICD-10 CODES THAT SUPPORT MEDICAL NECESSITY: Added: ICD-10 diagnosis codes E08.610, E09.610, E10.610, and E11.610 to Group 2 Codes for L4631 *06/04/2020: At this time 21st Century Cures Act applies to new and revised LCDs which require comment and notice. This revision is to an article that is not a local coverage determination.*
01/01/2020	R8	Revision Effective Date: 01/01/2020 REQUIREMENTS FOR SPECIFIC DMEPOS ITEMS PURSUANT TO FINAL RULE 1713 (84 Fed. Reg Vol 217): Added: Section and related information based on Final Rule 1713 POLICY SPECIFIC DOCUMENTATION REQUIREMENTS: Revised: "ordering physician's" to "treating practitioner's" Revised: "physician's" to "practitioner's" Revised: Format of HCPCS code references, from code 'spans' to individually-listed HCPCS CODING GUIDELINES: Revised: Format of HCPCS code references, from code 'spans' to individually-listed HCPCS Added: Coding Guidelines for L1900, L1902, L1904, L1907, L1910, L1920, L1930, L1932, L1940, L1945, L1950, L1951, L1970, L1971, L1980, and L1990 Revised: L1906 Coding Guideline Revised: L2006 Coding Guideline per quarterly HCPCS code update Removed: HCPCS K0903 Added: HCPCS A5514, crosswalk from K0903 Removed: Reference of effective DOS for K0903 Added: Reference of effective DOS for A5514 *02/20/2020: At this time 21st Century Cures Act applies to new and revised LCDs which require comment and notice. This revision is to an article that is not a local coverage determination.*
01/01/2020	R7	Revision Effective Date: 01/01/2020 CODING GUIDELINES: Added: L2006 Coding Guideline ICD-10 CODES THAT SUPPORT MEDICAL NECESSITY: Revised: Section header "ICD-10 Codes that are Covered" updated to "ICD-10 Codes that Support Medical Necessity" ICD-10 CODES THAT DO NOT SUPPORT MEDICAL NECESSITY: Revised: Section header "ICD-10 Codes that are Not Covered" updated to "ICD-10 Codes that DO NOT Support Medical Necessity" *12/19/2019: At this time 21st Century Cures Act applies to new and revised LCDs which require comment and notice. This revision is to an article that is not*

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION
		a local coverage determination.
01/01/2019	R6	Revision Effective Date: 01/01/2019 CODING GUIDELINES: Clarified: Custom fit requirements Revised: Coding instructions for prefabricated orthoses without distinction of OTS or custom-fit. Revised: RT and LT modifier billing instructions (Effective 03/01/2019) ICD-10 CODES THAT ARE COVERED: Added: All diagnosis codes formerly listed in the LCD ICD-10 CODES THAT ARE NOT COVERED: Added: Notation excluding unlisted diagnosis codes from coverage for specific HCPCS codes. Notation that for all other HCPCS codes, diagnoses are not specified. *03/28/2019: At this time 21st Century Cures Act applies to new and revised LCDs which require comment and notice. This revision is to an article that is not a local coverage determination.*
01/01/2017	R5	Revision Effective Date: 01/01/2017 CODING GUIDELINES: Revised: Code pairs to accurately reflect parallel codes Updated: HCPCS code narratives to align with HCPCS code table Added: Walking boot add-on bundling information *04/05/2018: At this time 21st Century Cures Act applies to new and revised LCDs that restrict coverage, which require comment and notice. This revision is to an article that is not a local coverage determination.*
01/01/2017	R4	Revision Effective Date: 01/01/2017 NON-MEDICAL NECESSITY COVERAGE AND PAYMENT RULES: Revised: Brace Benefit explanation to remove reference to "counterforce" that is no longer applicable Revised: Prefabricated and off-the-shelf (OTS) "minimal self-adjustment" regulatory definition discussion to improve consistency with regulatory definition of minimal self-adjustment Deleted: A4466 Added: A4467 Added: Instructions for A9285 Added: Policy specific documentation requirements from LCD CODING GUIDELINES: Removed: Reference to classification algorithm summary Revised: OTS and custom-fit definitions to improve consistency with regulatory definition of "minimal self-adjustment"

REVISION HISTORY DATE	REVISION HISTORY NUMBER	REVISION HISTORY EXPLANATION
		Added: Section on coding of elastic and similar materials Deleted: A4466 Added: A4467 Added A9285 RELATED LOCAL COVERAGE DOCUMENTS: Added: LCD-related Standard Documentation Requirements Language Article
07/01/2016	R3	Effective July 1, 2016 oversight for DME MAC Articles is the responsibility of CGS Administrators, LLC 18003 and 17013 and Noridian Healthcare Solutions, LLC 19003 and 16013. No other changes have been made to the Articles.
01/01/2016	R2	**Revision Effective Date: 01/01/2016** CODING GUIDELINES: Added: L4361 "clerical correction"
10/01/2015	R1	Revision Effective Date: 01/01/2015 NON-MEDICAL NECESSITY COVERAGE AND PAYMENT RULES: Added: Information for hospital and SNF reimbursement CODING GUIDELINES: Added: Reference to classification algorithm summary

Associated Documents

Related Local Coverage Document(s)

Article(s)
A55426 - Standard Documentation Requirements for All Claims Submitted to DME MACs
LCD(s)
L33686 - Ankle-Foot/Knee-Ankle-Foot Orthosis

Related National Coverage Document(s)

N/A

Statutory Requirements URL(s)

N/A

Rules and Regulations URL(s)

N/A

CMS Manual Explanations URL(s)

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Other URL(s)

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