Local Coverage Determination (LCD)

Ankle-Foot/Knee-Ankle-Foot Orthosis

L33686

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Contractor Information
LCD Information

Document Information

LCD ID
L33686
LCD Title
Ankle-Foot/Knee-Ankle-Foot Orthosis
Proposed LCD in Comment Period
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AMA CPT / ADA CDT / AHA NUBC Copyright Statement

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 no ambulatory 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

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Group 1

(133 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
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

Code	Description
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, 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

Code	Description
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, 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

Code	Description
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
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 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

Code	Description
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
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
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

Code	Description
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 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

Associated Documents

Attachments

N/A

Related Local Coverage Documents

Articles

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