

Health and Recovery Services Administration (HRSA)



Prosthetic and Orthotic Devices

Billing Instructions

Chapter 388-543 WAC

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About this publication

This publication supersedes all previous HRSA Prosthetic and Orthotic Devices Billing Instructions.

Published by the Health and Recovery Services Administration Washington State Department of Social and Health Services

Note: The effective date and publication date for any particular page of this document may be found at the bottom of the page.

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Important Contacts

A provider may use HRSA's toll-free lines for questions regarding its programs; however, HRSA's response is based solely on the information provided to the [HRSA] representative at the time of the call or inquiry, and in no way exempts a provider from following the rules and regulations that govern HRSA's programs. [WAC 388-502-0020(2)].

Where do I call for information on becoming a DSHS provider, submitting a change of address or ownership, or to ask questions about the status of a provider application?

Provider Enrollment Unit 866.545.0544

Where do I send my claims?

Hard Copy Claims:

Division of Program Support PO Box 9248 Olympia WA 98507-9248

Magnetic Tapes/Floppy Disks:

Division of Program Support Claims Control PO Box 45560 Olympia, WA 98504-5560

How do I request prior authorization?

All authorization issues, questions or comments should be addressed to:

Write/Call:

Division of Health Services Quality Support Quality Utilization Section Durable Medical Equipment PO Box 45506 Olympia, WA 98504-5506 800.292.8064 360.586.5299 fax

How do I request a Limitation Extension?

Write/Call:

Division of Health Services Quality Support Quality Utilization Section Durable Medical Equipment PO Box 45506 Olympia, WA 98504-5506 800.292.8064 360.586.5299 fax

Where do I address reimbursement issues, questions, or comments?

DME - Program Manager Professional Rates Section Division of Operational Support Services PO Box 45510 Olympia, WA 98504-5510

Where do I call if I have questions regarding electronic billing?

Write/call:

Electronic Billing Unit PO Box 45512 Olympia, WA 98504-5512 360.725.1267

Important Contacts (cont.)

How do I obtain copies of billing instructions or numbered memoranda?

Check out our web site at:

http://maa.dshs.wa.gov

Or write/call:

Provider Relations PO Box 45562 Olympia WA 98504-5562 800.562.3022

Who do I contact if I have questions regarding...

<u>Payments, denials, general questions</u> <u>regarding claims processing, or Healthy</u> <u>Options?</u>

Provider Relations 800.562.3022

Private insurance or third party liability, other than Healthy Options?

Coordination of Benefits Section 800.562.6136

Definitions & Abbreviations

This section defines terms and abbreviations (includes acronyms) used in these billing instructions.

Artificial limb – See prosthetic device. [WAC 388-543-1000]

By Report (BR) – A method of reimbursement for covered items, procedures, and services for which the department has no set maximum allowable fees. [WAC 388-543-1000]

Client - An applicant for, or recipient of, DSHS medical care program.

Code of Federal Regulations (CFR) - A codification of the general and permanent rules published in the federal register by the executive departments and agencies of the federal government.

Community Services Office (CSO) - An office of the department that administers social and health services at the community level. [WAC 388-500-0005]

Core Provider Agreement - The basic contract that HRSA holds with providers serving HRSA clients. The provider agreement outlines and defines terms of participation in Medical Assistance.

Date of Delivery – The date the client actually took physical possession of an item or equipment. [WAC 388-543-1000]

Department - The state Department of Social and Health Services (DSHS). [WAC 388-500-0005]

Expedited Prior Authorization – The process for obtaining authorization for selected durable medical equipment, and related supplies, prosthetics, orthotics, medical supplies and related services, in which providers use a set of numeric codes to indicate to HRSA which acceptable indications/conditions/HRSA-defined criteria are applicable to a particular request for DME authorization.

[WAC 388-543-1000]

Explanation of Benefits (EOB) - A coded message on the Medical Assistance Remittance Advice and Status Report (RA) that gives detailed information about the claim associated with that report.

Explanation of Medicare Benefits (**EOMB**) – A federal report generated for Medicare providers displaying transaction information regarding Medicare claims processing and payments.

Fee-for-Service – The general payment method HRSA uses to reimburse for covered medical services provided to clients, except those services covered under HRSA's prepaid managed care programs.

[WAC 388-543-1000]

Prosthetic and Orthotic Devices

Health and Recovery Services
Administration (HRSA) - The
administration within DSHS authorized by
the secretary to administer the acute care
portion of Title XIX Medicaid, Title XXI
State Children's Health Insurance Program
(SCHIP), Title XVI Supplemental Security
Income for the Aged, Blind, and Disabled
(SSI), and the state-funded medical care
programs, with the exception of certain
nonmedical services for persons with
chronic disabilities.

Health Care Financing Administration Common Procedure Coding System (HCPCS) – A coding system established by the Health Care Financing Administration to define services and procedures. [WAC 388-543-1000]

Internal Control Number (ICN) - A 17-digit number that appears on your Remittance Advice and Status Report (RA) by the client's name. Each claim is assigned an ICN when it is received by HRSA. The number identifies that claim throughout the claim's history.

Limitation Extension – A process for requesting and approving covered services and reimbursement that exceeds a coverage limitation (quantity, frequency, or duration) set in WAC, billing instructions, or numbered memoranda. Limitation extensions require prior authorization. [WAC 388-543-1000]

Managed Care - A prepaid comprehensive system of medical and health care delivery including preventive, primary, specialty, and ancillary health services.
[WAC 388-538-050]

Maximum Allowable - The maximum dollar amount for which a provider may be reimbursed by HRSA for specific services, supplies, or equipment.

Medicaid - The state and federally funded aid program that covers the Categorically Needy (CNP) and Medically Needy (MNP) programs.

Medical Identification card(s) – Medical Identification cards are the forms DSHS uses to identify clients of medical programs. These cards are good only for the dates printed on them. Clients will receive a Medical Identification card in the mail each month they are eligible. These cards are also known as DSHS Medical ID cards and were formerly called medical coupons or MAID cards.

Medical Management, Division of (DMM)

- A division within the Medical Assistance Administration responsible for the administration of the quality improvement and assurance programs, utilization review and management, and prior authorization for fee-for-service program.

Medically Necessary - A term for describing [a] requested service which is reasonably calculated to prevent, diagnose, correct, cure, alleviate or prevent worsening of conditions in the client that endanger life, or cause suffering or pain, or result in an illness or infirmity, or threaten to cause or aggravate a handicap, or cause physical deformity or malfunction. There is no other equally effective, more conservative or substantially less costly course of treatment available or suitable for the client requesting the service. For the purpose of this section, "course of treatment" may include mere observation or, where appropriate, no treatment at all. [WAC 388-500-0005]

Prosthetic and Orthotic Devices

Medicare - The federal government health insurance program for certain aged or disabled clients under Titles II and XVIII of the Social Security Act. Medicare has two parts:

- "Part A" covers the Medicare inpatient hospital, post-hospital skilled nursing facility care, home health services, and hospice care.
- "Part B" is the supplementary medical insurance benefit (SMIB) covering the Medicare doctor's services, outpatient hospital care, outpatient physical therapy and speech pathology services, home health care, and other health services and supplies not covered under Part A of Medicare. [WAC 388-500-0005]

Orthotic Device or Orthotic – A corrective or supportive device that:

- Prevents or corrects physical deformity or malfunction; or
- Supports a weak or deformed portion of the body. [WAC 388-543-1000]

Patient Identification Code (PIC) - An alphanumeric code that is assigned to each HRSA client consisting of:

- First and middle initials (a dash (-) must be entered if the middle initial is not indicated).
- Six-digit birthdate, consisting of numerals only (MMDDYY).
- First five letters of the last name (and spaces if the name is fewer than five letters).
- Alpha or numeric character (tiebreaker).

Prior Authorization – A process by which clients or providers must request and receive HRSA approval for certain medical equipment and related supplies, prosthetics, orthotics, medical supplies and related services, based on medical necessity, before the services are provided to clients, as a precondition for provider reimbursement. Expedited prior authorization and limitation extension are types of prior authorization. Also see WAC 388-501-0165. [WAC 388-543-1000]

Program Support, Division of (DPS) – The division within HRSA responsible for providing administrative services for the following:

- Claims Processing;
- Family Planning Services;
- Administrative Match Services to Schools and Health Departments;
- Managed Care Contracts; and
- Provider Enrollment/Relations

Prosthetic device or prosthetic – A replacement, corrective, or supportive device prescribed by a physician or other licensed practitioner of the healing arts, within the scope of his or her practice as defined by state law, to:

- Artificially replace a missing portion of the body;
- Prevent or correct physical deformity or malfunction; or
- Support a weak or deformed portion of the body. [WAC 388-543-1000]

Provider or Provider of Service - An institution, agency, or person:

- Who has a signed agreement [Core Provider] with the department to furnish medical care, goods, and/or services to clients; and
- Is eligible to receive payment from the department. [WAC 388-500-0005]

Remittance Advice and Status Report (Referred to as "RAs")- A report produced by the Medicaid Management Information System (MMIS) that provides detailed information concerning submitted claims and other financial transactions.

Resource Based Relative Value Scale (RBRVS) – A scale that measures the relative value of a medical service or intervention, based on amount of physician resources involved. [WAC 388-543-1000]

Revised Code Of Washington (RCW) - Washington State laws.

Third Party - Any entity that is or may be liable to pay all or part of the medical cost of care of a federal Medicaid or state medical program client.

Usual and Customary Charge – The amount the provider typically charges to 50% or more of his or her non-Medicaid clients, including clients with other third-party coverage. [WAC 388-543-1000]

Washington Administrative Code (WAC) - Codified rules of the State of Washington.

About the Program

What is the purpose of the Prosthetic and Orthotic Devices program? (Refer to WAC 388-543-1100)

The Medical Assistance Administration's (HRSA) Prosthetic and Orthotic (P&O) Devices program makes the purchase of medically necessary P&O devices accessible to eligible HRSA clients when the P&O devices are not included in other reimbursement methodologies (e.g., inpatient hospital DRG, nursing facility daily rate, HMO, or managed health care programs). The federal government deems P&O devices as optional services under the Medicaid program, except when:

- Prescribed as an integral part of an approved plan of treatment under the Home Health program; or
- Required under the Early and Periodic Screening, Diagnosis and Treatment (EPSDT) program.

Notifying Clients of Their Rights (Advance Directives) (42 CFR, Subpart I)

All Medicare-Medicaid certified hospitals, nursing facilities, home health agencies, personal care service agencies, hospices, and managed health care organizations are federally mandated to give **all adult clients** written information about their right, under state law, to make their own health care decisions.

Clients have the right to:

- Accept or refuse medical treatment;
- Make decisions concerning their own medical care; and
- Formulate an advance directive, such as a living will or durable power of attorney, for their health care.

| | | | Prosthetic and Or | thatic Davices |
|-------------|---------------|---------------|--------------------|----------------|
| | | | Trostilette and Or | inotic Devices |
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Client Eligibility

Who is eligible for P&O Devices? (Refer to Chapter 388-529 WAC)

Clients presenting Medical Identification cards with the following identifiers* **are eligible** for P&O devices:

| Medical Program Identifier | Medical Program |
|------------------------------|--|
| CNP | Categorically Needy Program – These clients are dual eligible (Medicare/Medicaid) |
| CNP Children's Health | Categorically Needy Program - Children's Health |
| CNP CHIP | Categorically Needy Program - Children's Health Insurance Program |
| GA-U No Out of State Care | General Assistance - Unemployable |
| LCP MNP | Limited Casualty Program-Medically Needy Program |
| MNP QMB | Medically Needy Program-Qualified Medicare Beneficiaries – These clients are dual eligible (Medicare/Medicaid) |

*Note: To clarify, clients presenting Medical ID cards with the following identifiers are not eligible for P&O devices:

- ✓ **QMB-Medicare Only** (Qualified Medicare Beneficiary-Medicare Only) (See *Billing* section)
- ✓ **MIP-EMER Hospital Only No out-of-state care** (Medically Indigent Program-EMER Hospital Only No out-of-state care)

Are clients enrolled in Healthy Options managed care eligible for P&O devices? (Refer to WAC 388-538-060 and 095)

YES! Clients with an identifier in the HMO column on their Medical Identification card are enrolled in one of HRSA's Healthy Options managed care plans. All services must be requested directly through the client's Primary Care Provider (PCP). Clients can contact their PCP by calling the telephone number located on their Medical Identification card.

All medical services covered under a managed health care plan must be obtained by the client through designated facilities or providers. The managed care plan is responsible for:

- Payment of covered services; and
- Payment of services referred by a provider participating with the plan to an outside provider.

To prevent billing denials, please check the client's Medical Identification card **prior** to scheduling services and at the **time of service** to make sure proper authorization or referral is obtained from the PCP and/or plan.

HRSA does not cover P&O devices provided by a nonparticipating provider for a client who is enrolled in a HRSA-contracted managed care plan. (**Refer to WAC 388-543-1400** [9])

Are clients enrolled in Primary Care Case Manager/Management (PCCM) eligible for P&O devices?

Yes! For the client who has chosen to obtain care with a PCCM, the identifier in the HMO column will be "PCCM." These clients must obtain or be referred for services via the PCCM. The PCCM is responsible for coordination of care just like the PCP would be in a plan setting. Please refer to the client's Medical Identification card for the PCCM. (See the *Billing* section for further information.)

Note: To prevent billing denials, please check the client's Medical Identification card **prior** to scheduling services and at the **time of the service** to make sure proper authorization or referral is obtain from the PCCM.

Coverage

What is covered? (Refer to WAC 388-543-1100)

- The Department of Social and Health Services (DSHS) covers the P&O devices, repairs, and labor charges listed in the *Fee Schedule* (section H) of this billing instruction.
- DSHS covers a replacement prosthesis only when the purchase of a replacement prosthesis is less costly than repairing or modifying a client's current prosthesis. (WAC 388-543-2600[3])

Note: Those HCPCS codes with a "#" symbol in the maximum allowable column of the fee schedule are not covered by DSHS.

What are the general conditions of coverage?

(Refer to WAC 388-543-1100)

DSHS covers the P&O devices listed in the *Fee Schedule* (section H) of this billing instruction when all of the following apply. The P&O devices must be:

- Medically necessary (see *Definitions* section). The provider or client must submit sufficient objective evidence to establish medical necessity. Information used to establish medical necessity includes, but is not limited to, the following:
 - A physiological description of the client's disease, injury, impairment, or other ailment, and any changes in the client's condition written by the prescribing physician, licensed prosthetist and/or orthotist, physical therapist, occupational therapist, or speech therapist; or
 - ✓ Video and/or photograph(s) of the client demonstrating the impairments and the client's ability to use the requested equipment, when applicable.
- Within the scope of an eligible client's medical care program (see *Client Eligibility* section);
- Within accepted medical or physical medicine community standards of practice;
- Prior authorized (see *Authorization* section);

- Prescribed by a physician or other licensed practitioner of the healing arts and within the scope of his or her practice as defined by state law. The prescription must state the specific item or service requested, diagnosis, prognosis, estimated length of need (weeks or months, not to exceed six months before being reevaluated), and quantity; and
- All written prior authorization requests must include a copy of the prescription upon submittal.
- Billed to the department as the payer of last resort only. DSHS does not pay first and then collect from Medicare.

Note: DSHS evaluates By Report (BR) items, procedures, or services for medical appropriateness and reimbursement value on a case-by-case basis.

What if a service is covered but considered experimental or has restrictions or limitations? (Refer to WAC 388-543-1100 [3] and [4])

- DSHS evaluates a request for a service that is in a covered category, but has been determined to be experimental or investigational as defined by WAC 388-531-0050, under the provisions of WAC 388-501-0165 which relate to medical necessity.
- DSHS evaluates a request for a covered service that is subject to limitations or other restrictions and approves such a service beyond those limitations or restrictions when medically necessary, under the standards for covered services in WAC 388-501-0165 (see page F.3 for limitation extensions).

How can I request that equipment/supplies be added to the "covered" list in this billing instruction? (WAC 388-543-1100 [7])

An interested party may request DSHS to include new P&O devices and related supplies and services in these billing instructions by sending a written request to DSHS's Quality Utilization Section (see *Important Contacts* section), plus all of the following:

- Manufacturer's literature;
- Manufacturer's pricing;
- Clinical research/case studies (including FDA approval, if required); and
- Any additional information the requestor feels is important.

What is not covered? (Refer to WAC 388-543-1300)

DSHS pays only for P&O devices and related supplies and services that are medically necessary, listed as covered, meet the definition of prosthetics and orthotics (see *Definitions* section), and prescribed per the provider requirements in this billing instruction (see *Provider Requirements* section).

DSHS considers all requests for covered P&O devices and related supplies and services, and noncovered P&O devices and related supplies and services, under the provisions of WAC 388-501-0165 which relate to medical necessity. When DSHS considers that a request does not meet the requirements for medical necessity, the definition(s) of covered item(s), or is not covered, the client may appeal that decision under the provisions of WAC 388-501-0165.

DSHS specifically excludes services and equipment in this billing instruction from fee-for-service (FFS) scope of coverage when the services and equipment do not meet the definition for a covered item, or the services are not typically medically necessary. This exclusion does not apply if the services and equipment are:

- Required under the EPSDT program;
- Included as part of a managed care plan service package;
- Included in a waivered program; or
- Part of one of the Medicare programs for qualified Medicare beneficiaries.

Services and equipment that are not covered include, but are not limited to:

- Services, procedures, devices, or the application of associated services that the Food and Drug Administration (FDA) and/or the Centers for Medicare and Medicaid Services (CMS) (formerly known as Health Care Financing Administration [HCFA]) consider investigative or experimental on the date the services are provided;
- Any service specifically excluded by statute;
- More costly services or equipment when DSHS determines that less costly, equally
 effective services or equipment are available;
- Hairpieces or wigs;
- Material or services covered under manufacturer's warranties:
- Procedures, prosthetics, or supplies related to gender dysphoria surgery;
- Shoe lifts less than one inch, arch supports, and nonorthopedic shoes;
- Supplies and equipment used during a physician office visit, such as tongue depressors and surgical gloves;

- Prosthetic devices dispensed for cosmetic reasons;
- Personal and comfort items that do not meet the definition of a prosthetic or orthotic device (see *Definitions* section), including, but not limited to, the following:
 - ✓ Clothing and accessories, such as coats, gloves (including wheelchair gloves), hats, scarves, slippers, and socks;
 - ✓ Cosmetics, including corrective formulations, hair depilatories, and products for skin bleaching, commercial sun screens, and tanning; and
 - ✓ Impotence devices;

Note: DSHS evaluates a request for any equipment or devices that are listed as noncovered in this billing instruction under the provisions of WAC 388-501-0165. (Refer to WAC 388-543-1100[2])

Prosthetic and Orthotic Devices Coverage Table

| Column | Abbreviation | Definition |
|---------------|--------------|--|
| Code Status | # | Non-covered item |
| Indicators | N | New |
| | D | Discontinued |
| | U | Update |
| | P | Policy Change |
| PA | Y | Requires Prior Authorization |
| PA | Y* | Requires Prior Authorization for clients 17 years of age and |
| | | older |
| Lic (License) | Y | Licensure required |
| Lic (License) | Y** | Licensure required if prescribing treatment of scoliosis |
| Lic (License) | *** | The item can be provided by a DME or Pharmacy provider as |
| | | long as other licensure requirements have been met |

| Code Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
|------------------------------|-------------------|----|-----|---|---------------------|
| # | A4280 | | | Adhesive skin support attachment for use with external breast prosthesis, each | |
| | A5500 | | | For diabetics only, fitting (including follow-up) custom preparation and supply of off-the-shelf depthinlay shoe manufactured to accommodate multidensity insert(s), per shoe | |
| | A5501 | | | For diabetics only, fitting (including follow-up) custom preparation and supply of shoe molded from cast(s) of patient's foot (custom molded shoe), per shoe | |
| | A5503 | | | For diabetics only, modification (including fitting) of off-the-shelf depth-inlay shoe or custom molded shoe with roller or rigid rocker bottom, per shoe | |
| | A5504 | | | For diabetics only, modification (including fitting) of off-the-shelf depth-inlay shoe or custom molded shoe with wedges, per shoe | |
| | A5505 | | | For diabetics only, modification (including fitting) of off-the-shelf depth-inlay shoe or custom molded shoe with metatarsal bar, per shoe | |
| | A5506 | | | For diabetics only, modification (including fitting) of off-the-shelf depth-inlay shoe or custom molded shoe with off-set heel(s), per shoe | |

| Code Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
|---|-------------------|----|-----|---|---------------------|
| marcators | A5507 | Y | Lic | For diabetics only, not otherwise specified | Comments |
| | A3307 | 1 | | modification (including fitting) of off-the-shelf depth- | |
| | | | | inlay or custom molded shoe, per shoe | |
| # | A5508 | | | For diabetics only, deluxe feature of off-the-shelf | |
| ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,, | 110000 | | | depth-inlay shoe or custom molded shoe, per shoe | |
| # | A5510 | | | For diabetics only, direct formed, compression molded | |
| | | | | to patient's foot without external heat source, multiple | |
| | | | | density insert(s) prefabricated, per shoe | |
| | A5512 | | | For diabetics only, multiple density insert, direct | |
| | | | | formed, molded to foot after external heat source of | |
| | | | | 230 degrees fahrenheit or higher, total contact with | |
| | | | | patient's foot, including arch, base layer minimum of | |
| | | | | 1/4 inch material of shore a 35 durometer or 3/16 inch | |
| | | | | material of shore a 40 durometer (or higher), | |
| | | | | prefabricated, each | |
| | A5513 | | | For diabetics only, multiple density insert, custom | |
| | | | | molded from model of patient's foot, total contact with | |
| | | | | patient's foot, including arch, base layer minimum of 3/16 inch material of shore a 35 durometer or higher, | |
| | | | | includes arch filler and other shaping material, custom | |
| | | | | fabricated, each | |
| # | E1800 | | | Dynamic adjustable elbow extension/flexion device, | |
| ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,, | 21000 | | | includes soft interface material | |
| # | E1801 | | | Static progressive stretch elbow device, extension | |
| | | | | and/or flexion, with or without range of motion | |
| | | | | adjustment, includes all components and accessories. | |
| # | E1802 | | | Dynamic adjustable forearm pronation/supination | |
| | | | | device, includes soft interface material | |
| # | E1805 | | | Dynamic adjustable wrist extension/flexion device, | |
| | | | | includes soft interface material | |
| # | E1806 | | | Static progressive stretch wrist device, flexion and/or | |
| | | | | extension, with or without range of motion | |
| ., | F1010 | | | adjustment, includes all components and accessories. | |
| # | E1810 | | | Dynamic adjustable knee extension/flexion device, | |
| ш | E1011 | | | includes soft interface material | |
| # | E1811 | | | Static progressive stretch knee device, flexion and/or extension, with or without range of motion | |
| | | | | adjustment, includes all components and accessories. | |
| # | E1815 | | | Dynamic adjustable ankle extension/flexion, includes | |
| π | L1013 | | | soft interface material | |
| # | E1816 | | | Static progressive stretch ankle device, flexion and/or | |
| " | 21010 | | | extension, with or without range of motion | |
| | | | | adjustment, includes all components and accessories. | |
| # | E1818 | | | Bi-directional progressive stretch forearm | |
| | | | | pronation/supination device with range of motion | |
| | | | | adjustment, includes cuffs | |

| Code Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
|------------------------------|-------------------|-----|-----|---|-------------------------|
| # | E1820 | 111 | Lic | Replacement soft interface material, dynamic | Comments |
| | | | | adjustable extension/flexion device | |
| # | E1821 | | | Replacement soft interface material/cuffs for bi- directional static progressive stretch device | |
| # | E1825 | | | Dynamic adjustable finger extension/flexion device, includes soft interface material | |
| # | E1830 | | | Dynamic adjustable toe extension/flexion device, includes soft interface material | |
| # | E1840 | | | Dynamic adjustable shoulder flexion/abduction/rotation device, includes soft interface material | |
| # | E1841 | | | Static progressive stretch shoulder device, with or without range of motion adjustment, includes all components and accessories. | |
| | K0672 | Y | | Addition to lower extremity orthosis, removable soft interface, all components, replacement only, each. | |
| | L0112 | Y | Y | Cranial cervical orthosis, congenital torticollis type, with or without soft interface material, adjustable range of motion joint, custom fabricated | |
| | L0113 | Y | Y | Cranial cervical orthosis, torticollis type, with or without joint, with or without soft interface material, prefabricated, includes fitting and adjustment | |
| | L0120 | | *** | Cervical, flexible, nonadjustable (foam collar) | |
| | L0130 | | Y | Cervical, flexible, thermoplastic collar, molded to patient | |
| | L0140 | | *** | Cervical, semi-rigid, adjustable (plastic collar) | |
| | L0150 | | *** | Cervical, semi-rigid, adjustable molded chin cup (plastic collar with mandibular/occipital piece) | |
| | L0160 | | | Cervical, semi-rigid, wire frame occipital/mandibular support | |
| | L0170 | Y | Y | Cervical, collar, molded to patient model | |
| | L0172 | - | *** | Cervical, collar, semi-rigid thermoplastic foam, two piece | |
| | L0174 | | *** | Cervical, collar, semi-rigid, thermoplastic foam, two piece with thoracic extension | |
| | L0180 | | | Cervical, multiple post collar, occipital/mandibular supports, adjustable | |
| | L0190 | | | Cervical, multiple post collar, occipital/mandibular supports, adjustable cervical bars (Somi, Guilford, Taylor types) | |
| | L0200 | | | Cervical, multiple post collar, occipital/mandibular supports, adjustable cervical bars, and thoracic extension | |
| D | L0210 | - | *** | Thoracic, rib belt | Removed January 2010 |

| Code | | | | | |
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| | Procedure | | T to | Description | Policy/ |
| Indicators | Code L0220 | PA | Lic *** | Description Thoracic, rib belt, custom fabricated | Comments |
| | L0220 | Y | Y** | Spinal orthosis, anterior-posterior-lateral control, with | |
| | L0430 | 1 | 1 | interface material, custom fitted (dewall posture | |
| | | | | protector only) | |
| | L0450 | | Y** | TLSO, flexible, provides trunk support, upper thoracic | |
| | | | | region, produces intracavitary pressure to reduce load | |
| | | | | on the intevertebral disks with rigid stays or panel(s), | |
| | | | | includes shoulder straps and closures, prefabricated, | |
| | 70470 | | | includes fitting and adjustment | |
| | L0452 | Y | | TLSO, flexible, provides trunk support, upper thoracic | |
| | | | | region, produces intracavitary pressure to reduce load on the intevertebral disks with rigid stays or panel(s), | |
| | | | | includes shoulder straps and closures, custom | |
| | | | | fabricated | |
| | L0454 | | Y** | TLSO, flexible, provides trunk support, extends from | |
| | | | _ | sacrococcygeal junction to above T-9 vertebra, | |
| | | | | restricts gross truck motion in the sagittal plane, | |
| | | | | produces intracavitary pressure to reduce load on the | |
| | | | | intervertebral disks with rigid stays or panel(s), | |
| | | | | includes shoulder straps and closures, prefabricated, | |
| | | | | includes fitting and adjustment | |
| | L0456 | | Y** | TLSO, flexible, provides trunk support, thoracic region, | |
| | | | | rigid posterior panel and soft anterior apron, extends | |
| | | | | from the sacrococcygeal junction and terminates just | |
| | | | | inferior to the scapular spine, restricts gross truck | |
| | | | | motion in the sagittal plane, produces intracavitary pressure to reduce load on the intervertebral disks, | |
| | | | | includes straps and closures, prefabricated, includes | |
| | | | | fitting and adjustment | |
| | L0458 | | Y | TLSO, triplanar control, modular segmented spinal | |
| | | | | system, two rigid plastic shells, posterior extends from | |
| | | | | the sacrococcygeal junction and terminates just | |
| | | | | inferior to the scapular spine, anterior extends from | |
| | | | | the symphysis pubis to the xiphold, soft liner, restricts | |
| | | | | gross trunk motion in the sagittal, coronal, and | |
| | | | | tranverse planes, lateral strength is provided by | |
| | | | | overlapping plastic and stabilizing closures, includes | |
| | | | | straps and closures, prefabricated, including fitting and adjustment | |
| | | | | and adjustinent | |

| Code Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
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| | L0460 | | Y | TLSO, triplanar control, modular segmented spinal system, two rigid plastic shells, posterior extends from the sacrococcygeal junction and terminates just inferior to the scapular spine, anterior extends from the symphysis pubis to the sternal notch, soft liner, restricts gross trunk motion in the sagittal, coronal, and tranverse planes, lateral strength is provided by overlapping plastic and stabilizing closures, includes straps and closures, prefabricated, including fitting and adjustment | Comments |
| | L0462 | | Y | TLSO, triplanar control, modular segmented spinal system, three rigid plastic shells, posterior extends from the sacrococcygeal junction and terminates just inferior to the scapular spine, anterior extends from the symphysis pubis to the sternal notch, soft liner, restricts gross trunk motion in the sagittal, coronal, and transverse planes, lateral strength is provided by overlapping plastic and stabilizing closures, includes straps and closures, prefabricated, including fitting and adjustment | |
| | L0464 | | Y | TLSO, triplanar control, modular segmented spinal system, four rigid plastic shells, posterior extends from the sacrococcygeal junction and terminates just inferior to scapular spine, anterior extends from symphysis pubis to the sternal notch, soft liner, restricts gross trunk motion in sagittal, coronal, and tranverse planes, lateral strength is provided by overlapping plastic and stabilizing closures, includes straps and closures, prefabricated, includes fitting and adjustment | |
| | L0466 | | Y | TLSO, sagittal-coronal control, rigid posterior frame and flexible soft anterior apron with straps, closures and padding, restricts gross trunk motion in sagittal plane, produces intracavitary pressure to reduce load on intervertebral disks, includes fitting and shaping the frame, prefabricated, includes fitting and adjustment | |
| | L0468 | | Y** | TLSO, sagittal control, rigid posterior frame and flexible soft anterior apron with straps, closures and padding, extends from sacroccoccygeal junction over scapulae, lateral strength provided by pelvic, thoracic and lateral frame pieces, restricts gross trunk motion in sagittal, and coronal planes, produces intracavitary pressure to reduce load on intervertebral disks, includes fitting and shaping the frame, prefabricated, includes filling and adjustment | |

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| Indicators | | PA | Lic | Description | Comments |
| | L0470 | | Y** | TLSO, triplanar-control, rigid posterior frame and | |
| | | | | flexible soft anterior apron with straps, closures and | |
| | | | | padding, extends from sacrococcygeal junction to | |
| | | | | scapula, lateral strength provided by pelvic, thoracic, | |
| | | | | and lateral frame pieces, rotational strength provided | |
| | | | | by subclavicular extentions, restricts gross trunk motion in sagittal, coronal, and tranverse planes, | |
| | | | | produces intracavitary pressure to reduce the load on | |
| | | | | intervertebral disks, includes fitting and shaping the | |
| | | | | frame, prefabricated, includes fitting and adjustment | |
| | L0472 | | Y** | TLSO, triplanar control, hyperextension, rigid anterior | |
| | L04/2 | | 1 | and lateral frame extends from symphysis pubis to | |
| | | | | sternal notch with two anterior components (one pubic | |
| | | | | and one sternal), posterior and lateral pads with straps | |
| | | | | and closures, limits spinal flexion, restricts gross trunk | |
| | | | | motion in sagittal, coronal, and transverse planes, | |
| | | | | includes fitting and shaping the frame, prefabricated, | |
| | | | | includes fitting and adjustment | |
| | L0474 | Y | Y** | TLSO, sagittal-coronal control, flexion compression | |
| | _== | _ | _ | jacket, two rigid plastic shells with soft liner, posterior | |
| | | | | extends from sacrococcygeal junction and terminates | |
| | | | | at or before the T-9 vertebra, anterior extends from | |
| | | | | symphysis pubis to xiphoid, usually laced together on | |
| | | | | one side, restricts gross trunk motion in sagittal and | |
| | | | | coronal planes, allows free flexion and compression of | |
| | | | | the LS region, includes straps and closures, | |
| | | | | prefabricated, includes fitting and adjustment | |
| | L0480 | Y | Y | TLSO, triplanar control, one piece rigid plastic shell | |
| | | | | without interface liner, with multiple straps and | |
| | | | | closures, posterior extends from sacrococcygeal | |
| | | | | junction and terminates just inferior to scapular spine, | |
| | | | | anterior extends from symphysis pubis to sternal | |
| | | | | notch, anterior or posterior opening, restricts gross | |
| | | | | trunk motion in sagittal, coronal, and transverse | |
| | | | | planes, includes a carved plaster or CAD-CAM model, | |
| | T 0 100 | • | ** | custom fabricated | |
| | L0482 | Y | Y | TLSO, triplanar control, one piece rigid plastic shell | |
| | | | | with interface liner, with multiple straps and closures, | |
| | | | | posterior extends from sacrococcygeal junction and | |
| | | | | terminates just inferior to scapular spine, anterior | |
| | | | | extends from symphysis pubis to sternal notch, | |
| | | | | anterior or posterior opening, restricts gross trunk | |
| | | | | motion in sagittal, coronal, and transverse planes, includes a carved plaster or CAD-CAM model, | |
| | | | | custom fabricated | |
| | | | | custom ratificated | |
| | | | | | |

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| Status Indicators | Procedure Code | PA | Lic | Dogavintian | Policy/ Comments |
| muicators | L0484 | Y | Y | TLSO, triplanar control, two piece rigid plastic shell | Comments |
| | L0464 | 1 | 1 | without interface liner, with multiple straps and | |
| | | | | closures, posterior extends from sacrococcygeal | |
| | | | | junction and terminates just inferior to scapular spine, | |
| | | | | anterior extends from symphysis pubis to sternal | |
| | | | | notch, lateral strength is enhanced by overlapping | |
| | | | | plastic, restricts gross trunk motion in sagittal, | |
| | | | | coronal, and transverse planes, includes a carved | |
| | I 0406 | 37 | *7 | plaster or CAD-CAM model, custom fabricated | |
| | L0486 | Y | Y | TLSO, triplanar control, two piece rigid plastic shell | |
| | | | | with interface liner, with multiple straps and closures, posterior extends from sacrococcygeal junction and | |
| | | | | terminates just inferior to scapular spine, anterior | |
| | | | | extends from symphysis pubis to sternal notch, lateral | |
| | | | | strength is enhanced by overlapping plastic, restricts | |
| | | | | gross trunk motion in sagittal, coronal, and transverse | |
| | | | | planes, includes a carved plaster or CAD-CAM model, | |
| | | | | custom fabricated | |
| | L0490 | | Y** | TLSO, sagittal-coronal control, one piece rigid plastic | |
| | | | | shell with overlapping reinforced anterior, with | |
| | | | | multiple straps and closures, posterior extends from sacrococcygeal junction and terminates at or before | |
| | | | | the T-9 vertebra, anterior extends from symphysis | |
| | | | | pubis to xiphoid, anterior opening, restricts gross trunk | |
| | | | | motion in sagittal and coronal planes, prefabricated, | |
| | | | | includes fitting and adjustment | |
| | L0491 | | Y | TLSO, Sagittal-coronal control, modular segmented | |
| | | | | spinal system, two rigid plastic shells, posterior | |
| | 1.0402 | | 3.7 | extends from the sacrococcygeal junction area | |
| | L0492 | | Y | TLSO, Sagittal-coronal control, modular segmented spinal system, three rigid plastic shells, posterior | |
| | | | | extends from the sacrococcygeal junction | |
| | L0621 | | Y** | Sacroiliac orthosis, flexible, provides pelvic-sacral | |
| | | | | support, reduces motion about the sacroiliac joint, | |
| | | | | includes straps, closures, may include pendulous | |
| | | | | abdomen design, prefabricated, includes fitting and | |
| | | | | adjustment | |
| | L0622 | | Y**/ *** | Sacroiliac orthosis, flexible, provides pelvic-sacral | |
| | | | ベベボ | support, reduces motion about the sacroiliac joint, | |
| | | | | includes straps, closures, may included pendulous abdomen design, custom fabricated | |
| | L0623 | Y | Y**/ | Sacroiliac orthosis, provides pelvic-sacral support, | |
| | L0023 | 1 | *** | with rigid or semi-rigid panels over the sacrum and | |
| | | | | abdomen, reduces motion about the sacroiliac joint, | |
| | | | | includes straps, closures, may include pendulous | |
| | | | | abdomen design, prefabricated, includes fitting and | |
| | | | | adjustment | |

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| Indicators | | PA | | Description | Comments |
| | L0624 | Y | Y** | Sacroiliac orthosis, provides pelvic-sacral support, with rigid or semi-rigid panels over the sacrum and abdomen, reduces motion about the sacroiliac joint, includes straps, closures, may include pendulous abdomen design, custom fabricated | |
| | L0625 | | Y**/ *** | Lumbar orthosis, flexible, provides lumbar support, posterior extends from L-1 to below L - 5 vertebra, produces intracavitary pressure to reduce load on the intervertebral discs, includes straps, closures, may include pendulous abdomen design, shoulder straps, stays, prefabricated, includes fitting and adjustment | |
| | L0626 | | Y**/ *** | Lumbar orthosis, sagittal control, with rigid posterior panel(s), posterior extends from L-1 to below L-5 vertebra, produces intracavitary pressure to reduce load on the intervertebral discs, includes straps, closures, may include padding, stays, shoulder straps, pendulous abdomen design, prefabricated, includes fitting and adjustment | |
| | L0627 | | / *** | Lumbar orthosis, sagittal control, with rigid anterior and posterior panels, posterior extends from L-1 to below L-5 vertebra, produces intractivitary pressure to reduce load on the intervertebral discs, includes straps, closures, may include padding, shoulder straps, pendulous abdomen design, prefabricated, includes fitting and adjustment | |
| | L0628 | | Y**/* ** | Lumbar-sacral orthosis, flexible, provides lumbo- sacral support, posterior extends from sacrococcygeal junction to T-9 vertebra, produces intracavitary pressure to reduce load on the intervertebral discs, includes straps, closures, may include stays, shoulder straps, pendulous abdomen design, prefabricated, includes fitting and adjustment | |
| | L0629 | Y | Y**/ *** | Lumbar-sacral orthosis, flexible, provides lumbo- sacral support, posterior extends from sacrococcygeal junction to T-9 vertebra, produces intracavitary pressure to reduce load on the intervertebral discs, includes straps, closures, may include stays, shoulder straps, pendulous abdomen design, custom fabricated | |
| | L0630 | | Y**/ *** | Lumbar-sacral orthosis, sagittal control, with rigid posterior panel(s), posterior extends from sacrococcygeal junction to T-9 vertebra, produces intracavitary pressure to reduce load on the intervertebral discs, includes straps, closures, may include padding, stays, shoulder straps, pendulous abdomen design, prefabricated, includes fitting and adjustment | |

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| | L0631 | | Y** | Lumbar-sacral orthosis, sagittal-coronal control, with rigid anterior and posterior panels, posterior extends from sacrococcygeal junction to T-9 vertebra, produces intracavitary pressure to reduce load on the intervertebral discs, includes straps, pendulous abdomen design, prefabricated, includes fitting and adjustment | |
| | L0632 | Y | Y | Lumbar-sacral orthosis, sagittal-coronal control, with rigid anterior and posterior panels, posterior extends from sacrococcygeal junction to T-9 vertebra, produces intracavitary pressure to reduce load on the intervertebral discs, includes straps, closures, may include padding, shoulder straps, pendulous abdomen design, custom fabricated | |
| | L0633 | | Y**/* ** | Lumbar-sacral orthosis, sagittal-coronal control, with rigid posterior frame/Panel(s), posterior extends from sacrococcygeal junction to T-9 vertebra, lateral strength provided by rigid lateral frame/panels, produces intracavitary pressure to reduce load on intervertebral discs, includes straps, closures, may include padding, stays, shoulder straps, pendulous abdomen design, prefabricated, includes fitting and adjustment | |
| | L0634 | Y | Y | Lumbar-sacral orthosis, sagittal-coronal control, with rigid posterior frame/Panel(s), posterior extends from sacrococcygeal junction to T-9 vertebra, lateral strength provided by rigid lateral frame/panels, produces intracavitary pressure to reduce load on intervertebral discs, includes straps, closures, may include padding, stays, shoulder straps, pendulous abdomen design, custom fabricated | |
| | L0635 | Y | Y | Lumbar-sacral orthosis, sagittal-coronal control, lumbar flexion, rigid posterior frame/panels, lateral articulating design to flex the lumbar spine, posterior extends from sacrococcygeal junction to T-9 vertebra, lateral strength provided by rigid lateral frame/panels, produces intracavitary pressure to reduce load on intervertebral discs, includes straps, closures, may include padding, anterior panel, pendulous abdomen design, prefabricated, includes fitting and adjustment | |

| | Procedure | | T ia | Decomination | Policy/ |
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| Indicators | | PA | Lic | Description | Comments |
| | L0636 | Y | Y | Lumbar-sacral orthosis, sagittal-coronal control, | |
| | | | | lumbar flexion, rigid posterior frame/panels, lateral | |
| | | | | articulating design to flex the lumbar spine, posterior | |
| | | | | extends from sacrococcygeal junction to T-9 vertebra, | |
| | | | | lateral strength provided by rigid lateral frame/panels, | |
| | | | | produces intracavitary pressure to reduce load on intervertebral discs, includes straps, closures, may | |
| | | | | include padding, anterior panel, pendulous abdomen | |
| | | | | design, custom fabricated | |
| | L0637 | Y | Y** | Lumbar-sacral orthosis, sagittal-coronal control, with | |
| | L0037 | 1 | 1 | rigid anterior and posterior frame/panels, posterior | |
| | | | | extends from sacrococcygeal junction to T-9 vertebra, | |
| | | | | lateral strength provided by rigid lateral frame/panels, | |
| | | | | produces intracavitary pressure to reduce load on | |
| | | | | intervertebral discs, includes straps, closures, may | |
| | | | | include padding, shoulder straps, pendulous abdomen | |
| | | | | design, prefabricated, includes fitting and adjustment | |
| | L0638 | Y | Y | Lumbar-sacral orthosis, sagittal-coronal control, with | |
| | | | | rigid anterior and posterior frame/panels, posterior | |
| | | | | extends from sacrococcygeal junction to T-9 vertebra, | |
| | | | | lateral strength provided by rigid lateral frame/panels, | |
| | | | | produces intracavitary pressure to reduce load on | |
| | | | | intervertebral discs, includes straps, closures, may | |
| | | | | include padding, shoulder straps, pendulous abdomen | |
| | | | | design, custom fabricated | |
| | L0639 | | Y^{**} | Lumbar-sacral orthosis, sagittal-control, rigid | |
| | | | | shell(s)/panel(s) posterior extends from | |
| | | | | sacrococcygeal junction to T-9 vertebra, anterior | |
| | | | | extends from symphysis pubis to xiphoid, produces | |
| | | | | intracavitary pressure to reduce load on the | |
| | | | | intervertebral discs, overall strength is provided by | |
| | | | | overlapping rigid material and stabilizing closures, | |
| | | | | includes straps, closures, may include soft interface, pendulous abdomen design, prefabricated, includes | |
| | | | | fitting and adjustment | |
| | L0640 | Y | Y** | Lumbar-sacral orthosis, sagittal-control, rigid | |
| | L0040 | 1 | 1 | shell(s)/panel(s) posterior extends from | |
| | | | | sacrococcygeal junction to T-9 vertebra, anterior | |
| | | | | extends from symphysis pubis to xiphoid, produces | |
| | | | | intracavitary pressure to reduce load on the | |
| | | | | intervertebral discs, overall strength is provided by | |
| | | | | overlapping rigid material and stabilizing closures, | |
| | | | | includes straps, closures, may include soft interface, | |
| | | | | pendulous abdomen design, custom fabricated | |
| | L0700 | Y | Y | CTLSO, anterior-posterior-lateral control, molded to | |
| | | | | patient model (Minerva type) | |

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| Indicators | Code | PA | Lic | Description | Comments |
| | L0710 | Y | Y | CTLSO, anterior-posterior-lateral control, molded to patient model, with interface material, (Minerva type) | |
| | L0810 | | Y | Halo procedure, cervical halo incorporated into jacket | |
| | L0820 | | Y | vest Halo procedure, cervical halo incorporated into plaster | |
| | L0830 | Y | Y | body jacket Halo procedure, cervical halo incorporated into | |
| | L0630 | 1 | 1 | Milwaukee type orthosis | |
| | L0859 | | Y | Addition to halo procedures, magnetic resonance image compatible system | |
| | L0861 | Y | Y | Addition to halo procedure, replacement | |
| | ¥ 00=0 | | **** | liner/interface material | |
| | L0970 | | Y** *** | TLSO, corset front | |
| | L0972 | | Y** | LSO, corset front | |
| | L0974 | | Y** | TLSO, full corset | |
| | L0976 | | Y | LSO, full corset | |
| | | | **/** * | · | |
| | L0978 | | *** | Axillary crutch extension | |
| | L0980 | | *** | Peroneal straps, pair | |
| | L0982 | | *** | Stocking supporter grips, set of four (4) | |
| | L0984 | Y | *** | Protective body sock, each | |
| | L0999 | Y | | Addition to spinal orthosis, not otherwise specified | |
| | L1000 | Y* | Y | CTLSO (Milwaukee), inclusive of furnishing initial | |
| | | | | orthosis, including model | |
| | L1001 | Y | Y** | Cervical thoracic lumbar sacral orthosis, immobilizer, infant size, prefabricated, includes fitting and adjustment | |
| | L1005 | Y | Y | Tension based scoliosis orthosis and accessory pads, includes fitting and adjustment | |
| | L1010 | | Y | Addition to CTLSO or scoliosis orthosis, axilla sling | |
| | L1020 | | Y | Addition to CTLSO or scoliosis orthosis, kyphosis pad | |
| | L1025 | | Y | Addition to CTLSO or scoliosis orthosis, kyphosis pad, floating | |
| | L1030 | | Y | Addition to CTLSO or scoliosis orthosis, lumbar bolster pad | |
| | L1040 | | Y | Addition to CTLSO or scoliosis orthosis, lumbar or lumbar rib pad | |
| | L1050 | | Y | Addition to CTLSO or scoliosis orthosis, sternal pad | |
| | L1060 | | Y | Addition to CTLSO or scoliosis orthosis, thoracic pad | |
| | L1070 | | Y | Addition to CTLSO or scoliosis orthosis, trapezius sling | |
| | L1080 | | Y | Addition to CTLSO or scoliosis orthosis, outrigger | |

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| Indicators | | PA | Lic | Description | Comments |
| | L1085 | | Y | Addition to CTLSO or scoliosis orthosis, outrigger, | |
| | | | | bilateral with vertical extensions | |
| | L1090 | | Y | Addition to CTLSO or scoliosis orthosis, lumbar sling | |
| | L1100 | | Y | Addition to CTLSO or scoliosis orthosis, ring flange, plastic or leather | |
| | L1110 | Y* | Y | Addition to CTLSO or scoliosis orthosis, ring flange, plastic or leather, molded to patient model | |
| | L1120 | Y* | Y | Addition to CTLSO or scoliosis orthosis, cover for upright, each | |
| | L1200 | Y* | Y | TLSO, inclusive of furnishing initial orthosis only | |
| | L1210 | | Y | Addition to TLSO, (low profile), lateral thoracic extension | |
| | L1220 | | Y | Addition to TLSO, (low profile), anterior thoracic extension | |
| | L1230 | | Y | Addition to TLSO, (low profile), Milwaukee type superstructure | |
| | L1240 | | Y | Addition to TLSO, (low profile), lumbar derotation pad | |
| | L1250 | | Y | Addition to TLSO, (low profile), anterior ASIS pad | |
| | L1260 | | Y | Addition to TLSO, (low profile), anterior thoracic derotation pad | |
| | L1270 | | Y | Addition to TLSO, (low profile), abdominal pad | |
| | L1280 | | Y | Addition to TLSO, (low profile), rib gusset (elastic), each | |
| | L1290 | | Y | Addition to TLSO, (low profile), lateral trochanteric pad | |
| | L1300 | Y* | Y | Other scoliosis procedure, body jacket molded to patient model | |
| | L1310 | Y* | Y | Other scoliosis procedures, postoperative body jacket | |
| | L1499 | Y | Y | Spinal orthosis, not otherwise specified | |
| | L1500 | Y | | THKAO, mobility frame (Newington, Parapodium types) | |
| | L1510 | | | THKAO, standing frame; with or without tray accessories | Limit of one per client every 5 years. |
| | L1520 | Y | | THKAO, swivel walker | |
| | L1600 | | | HO, abduction control of hip joints, flexible, Frejka type, with cover, prefabricated, includes fitting and adjustment | |
| | L1610 | | | HO, abduction control of hip joints, flexible, (Frejka cover only), prefabricated, includes fitting and adjustment | |
| | L1620 | | | HO, abduction control of hip joints, flexible, (Pavlik Harness), prefabricated, includes fitting and adjustment | |

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| Indicators | | PA | Lic | Description | Policy/ Comments |
| | L1630 | | Y | HO, abduction control of hip joints, semi-flexible | |
| | | | | (Von Rosen type), prefabricated, includes fitting and | |
| | L1640 | | Y | adjustment HO, abduction control of hip joints, static, pelvic band | |
| | L1040 | | 1 | or spreader bar, thigh cuffs, custom fabricated | |
| | L1650 | | | HO, abduction control of hip joints, static, adjustable, | |
| | | | | (Ilfled type), prefabricated, includes fitting and | |
| | L1652 | | | adjustment Hip orthosis, bilateral thigh cuffs with adjustable | |
| | L1032 | | | abductor spreader bar, adult size, prefabricated, | |
| | | | | includes fitting and adjustment, any type | |
| | L1660 | | | HO, abduction control of hip joints, static, plastic, prefabricated, includes fitting and adjustment | |
| | L1680 | | Y | HO, abduction control of hip joints, dynamic, pelvic | |
| | | | | control, adjustable hip motion control, thigh cuffs (Rancho hip action type), custom fabricated | |
| | L1685 | | Y | HO, abduction control of hip joint, postoperative hip | |
| | | | _ | abduction type, custom fabricated | |
| | L1686 | | Y | HO, abduction control of hip joint, postoperative hip | |
| | | | | abduction type, prefabricated, includes fitting and adjustment | |
| | L1690 | Y | Y | Combination, bilateral, lumbo-sacral, hip, femur | |
| | | | | orthosis providing adduction and internal rotation | |
| | I 1700 | 37 | X 7 | control, prefabricated, includes fitting and adjustment | |
| | L1700 | Y | Y | Legg Perthes orthosis (Toronto type), custom fabricated | |
| | L1710 | Y | Y | Legg Perthes orthosis (Newington type), custom fabricated | |
| | L1720 | Y | Y | Legg Perthes orthosis, trilateral (Tachdijan type), custom fabricated | |
| | L1730 | | Y | Legg Perthes orthosis (Scottish Rite type), custom fabricated | |
| | L1755 | Y | Y | Legg Perthes orthosis (Patten bottom type), custom | |
| | | | | fabricated | |
| D | L1800 | - | <u>***</u> | KO, elastic with stays, prefabricated, includes fitting | Removed |
| | L1810 | | *** | And adjustment KO, elastic with joints, prefabricated, includes fitting | January 2010 |
| | 21010 | | | and adjustment | |
| D | L1815 | - | *** | J 1 | Removed |
| | I 1000 | | *** | pad(s), prefabricated, includes fitting and adjustment | January 2010 |
| | L1820 | | ጥ ጥ ጥ | KO, elastic with condylar pads and joints, prefabricated, includes fitting and adjustment | |
| D | L1825 | _ | *** | KO, elastic knee cap, prefabricated, includes fitting | Removed |
| | | | | and adjustment | January 2010 |
| | L1830 | | *** | KO, Immobilizer, canvas longitudinal, prefabricated, | |
| | | | | includes fitting and adjustment | |

| Code Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
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| ZII GICGIOIS | L1831 | 111 | Lic | Knee orthosis, locking knee joint(s), positional | |
| | | | | orthosis, prefabricated, includes fitting and adjustment | |
| | L1832 | | | KO, adjustable knee joints (Unicentric or polycentric), | |
| | | | | positional orthosis, rigid support, prefabricated, | |
| | L1834 | Y | Y | includes fitting and adjustment KO, without knee joints, rigid, custom fabricated | |
| | L1836 | 1 | 1 | Knee orthosis, rigid, without joint(s), includes soft | |
| | 21030 | | | interface material, prefabricated, includes fitting and adjustment | |
| | L1840 | | Y | KO, derotation, medial-lateral, anterior cruciate ligament, custom fabricated | |
| | L1843 | | | KO, single upright, thigh and calf, with adjustable flexion and extension joint (Unicentric or polycentric), medial-lateral and rotation control, prefabricated, includes fitting and adjustment | |
| | L1844 | Y | | KO, single upright, thigh and calf, with adjustable flexion and extension joint (Unicentric or polycentric), medial-lateral and rotation control, custom fabricated | |
| | L1845 | | | KO, double upright, thigh and calf, with adjustable flexion and extension joint (Unicentric or polycentric), medial-lateral and rotation control, prefabricated, includes fitting and adjustment | |
| | L1846 | Y | Y | KO, double upright, thigh and calf, with adjustable flexion and extension joint (Unicentric or polycentric), medial-lateral and rotation control, custom fabricated | |
| | L1847 | | | KO, double upright with adjustable joint, with inflatable air support chamber(s), prefabricated, includes fitting and adjustment | |
| | L1850 | | | KO, Swedish type, prefabricated, includes fitting and adjustment | |
| | L1860 | Y | Y | KO, modification of supracondylar prosthetic socket, custom fabricated (SK) | |
| | L1900 | | Y | AFO, spring wire, dorsiflexion assist calf band, custom fabricated | |
| D | L1901 | - | <u>***</u> | Ankle orthosis, elastic, prefabricated, includes fitting and adjustment (e.g., neoprene, Lycra) | Removed January 2010 |
| | L1902 | | *** | AFO, ankle gauntlet, prefabricated, includes fitting and adjustment | |
| | L1904 | | Y | AFO, molded ankle gauntlet, custom fabricated | |
| | L1906 | | *** | AFO, multiligamentus ankle support, prefabricated, includes fitting and adjustment | |
| | L1907 | | Y | AFO, supramalleolar with straps, with or without interface/pads, custom fabricated | |
| | L1910 | | | AFO, posterior, single bar, clasp attachment to shoe counter, prefabricated, includes fitting and adjustment | |

| | Procedure | | | | Policy/ |
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| Indicators | | PA | Lic | Description | Comments |
| | L1920 | | Y | AFO, 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 | | Y | Ankle foot orthosis, plastic or other material, custom fabricated | |
| | L1945 | Y | Y | AFO, molded to patient model, plastic, rigid anterior tibial section (floor reaction), custom fabricated | See EPA criteria, pages E.5-E.7. |
| | L1950 | Y | Y | AFO, spiral, (IRM type), plastic, custom fabricated | |
| | L1951 | Y | Y | Ankle foot orthosis, spiral, (institute of rehabilitative medicine type), plastic or other material, prefabricated, includes fitting and adjustment | |
| | L1960 | | Y | AFO, posterior solid ankle, plastic, custom fabricated | |
| | L1970 | | Y | AFO, plastic, with ankle joint, custom fabricated | |
| | L1971 | Y | Y | Ankle foot orthosis, plastic or other material with ankle joint, prefabricated, includes fitting and adjustment | |
| | L1980 | | Y | AFO, single upright free plantar dorsiflexion, solid stirrup, calf band/cuff (single bar "BK" orthosis), custom fabricated | |
| | L1990 | | Y | AFO, double upright free plantar dorsiflexion, solid stirrup, calf band/cuff (double bar "BK" orthosis), custom fabricated | |
| | L2000 | | Y | KAFO, single upright, free knee, free ankle, solid stirrup, thigh and calf bands/cuffs (single bar "AK" orthosis), custom fabricated | |
| | L2005 | Y | Y | Knee ankle foot orthosis, any material, single or double upright, stance control, automatic lock and swing phase release, mechanical activation, includes ankle joint, any type, custom fabricated | |
| | L2010 | | Y | KAFO, single upright, free ankle, solid stirrup, thigh and calf bands/cuffs (single bar "AK" orthosis), without knee joint, custom fabricated | |
| | L2020 | | Y | KAFO, double upright, free knee, free ankle, solid stirrup, thigh and calf bands/cuffs (double bar "AK" orthosis), custom fabricated | |
| | L2030 | | Y | KAFO, double upright, free ankle, solid stirrup, thigh and calf bands/cuffs (double bar "AK" orthosis), without knee joint, custom fabricated | |
| | L2034 | Y | Y | Knee ankle foot orthosis, full plastic, single upright, with or without free motion knee, medial lateral rotation control, with or without free motion | |

| | Procedure | | т. | | Policy/ | |
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| Indicators | | PA | Lic | Description VARO 6 II a circuit distribution | Comments | |
| | L2035 | | | KAFO, full plastic, static, (pediatric size), | | |
| | 1 2026 | 3.7 | 37 | prefabricated, includes fitting and adjustment | | |
| | L2036 | Y | Y | KAFO, full plastic, double upright, with or without | | |
| | | | | | free motion knee, with or without free motion ankle, | |
| | 1 2027 | 37 | 37 | custom fabricated | | |
| | L2037 | Y | Y | KAFO, full plastic, single upright, with or without free | | |
| | | | | motion knee, with or without free motion ankle, custom fabricated | | |
| | L2038 | Y | Y | KAFO, full plastic, with or without free motion knee, | | |
| | L2036 | 1 | 1 | with or without free motion ankle, multiaxis ankle, | | |
| | | | | (Lively orthosis or equal), custom fabricated | | |
| | L2040 | | Y | HKAFO, torsion control, bilateral rotation straps, | | |
| | L2040 | | 1 | pelvic band/belt, custom fabricated | | |
| | L2050 | | Y | HKAFO, torsion control, bilateral torsion cables, hip | | |
| | L2030 | | 1 | joint, pelvic band/belt, custom fabricated | | |
| | L2060 | | Y | HKAFO, torsion control, bilateral torsion cables, ball | | |
| | L2000 | | 1 | bearing hip joint, pelvic band/belt, custom fabricated | | |
| | L2070 | | Y | HKAFO, torsion control, unilateral rotation straps, | | |
| | L2070 | | 1 | pelvic band/belt, custom fabricated | | |
| | L2080 | | Y | HKAFO, torsion control, unilateral torsion cable, hip | | |
| | L2000 | | 1 | joint, pelvic band/belt, custom fabricated | | |
| | L2090 | | Y | HKAFO, torsion control, unilateral torsion cable, ball | | |
| | L2070 | | 1 | bearing hip joint, pelvic band/belt, custom fabricated | | |
| | L2106 | | Y | AFO, fracture orthosis, tibial fracture cast orthosis, | | |
| | 22100 | | • | thermoplastic type casting material, custom fabricated | | |
| | L2108 | Y | Y | AFO, fracture orthosis, tibial fracture cast orthosis, | | |
| | | | _ | custom fabricated | | |
| | L2112 | | | AFO, fracture orthosis, tibial fracture orthosis, soft, | | |
| | | | | prefabricated, includes fitting and adjustment | | |
| | L2114 | | | AFO, fracture orthosis, tibial fracture orthosis, semi- | | |
| | | | | rigid, prefabricated, includes fitting and adjustment | | |
| | L2116 | | | AFO, fracture orthosis, tibial fracture orthosis, rigid, | | |
| | | | | prefabricated, includes fitting and adjustment | | |
| | | | | | | |
| | L2126 | Y | Y | KAFO, fracture orthosis, femoral fracture cast | | |
| | | | | orthosis, thermoplastic type casting material, custom | | |
| | | | | fabricated | | |
| | L2128 | Y | Y | KAFO, 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 | | |

| | Procedure | D . | | D 1.4 | Policy/ |
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| Indicators | | PA | Lic | Description | Comments |
| | 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 | Y | Y | 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 | | Y | 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, nonmolded lacer | |

| | Procedure | | T:a | Description | Policy/ |
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| Indicators | | PA | Lic | Description Addition to be seen and additional to the seen additional to the seen additional to the seen and additional to the seen additio | Comments |
| | L2330 | | Y | Addition to lower extremity, lacer molded to patient model | |
| | L2335 | | | Addition to lower extremity, anterior swing band | |
| | L2333 | | Y | Addition to lower extremity, anterior swing band Addition to lower extremity, pretibial shell, molded to | |
| | | | | patient model | |
| | L2350 | | Y | 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 | Y | | 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 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 | | Y | Addition to lower extremity, thigh/weight bearing, quadric-lateral brim, molded to patient model | |
| | L2520 | | | Addition to lower extremity, thigh/weight bearing, quadric-lateral brim, custom fitted | |
| | L2525 | Y | Y | Addition to lower extremity, thigh/weight bearing, ischial containment/narrow M-L brim molded to patient model | |

| Code Status | Procedure | | | | Policy/ |
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| Indicators | | PA | Lic | Description | Comments |
| | 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, nonmolded | |
| | L2540 | | Y | Addition to lower extremity, thigh/weight bearing, | |
| | 1.2550 | | | lacer, molded to patient model | |
| | L2550 | | | Addition to lower extremity, thigh/weight bearing, high roll cuff | |
| | L2570 | | | Addition to lower extremity, pelvic control, hip joint | |
| | L2370 | | | Clevis type, two position joint, each | |
| | L2580 | | | Addition to lower extremity, pelvic control, pelvic | |
| | | | | sling | |
| | L2600 | | | Addition to lower extremity, pelvic control, hip joint, | |
| | | | | Clevis type, or thrust bearing, free, each | |
| | L2610 | | | Addition to lower extremity, pelvic control, hip joint, | |
| | 7.0.00 | | | Clevis or thrust bearing, lock, each | |
| | L2620 | | | Addition to lower extremity, pelvic control, hip joint, | |
| | L2622 | | | heavy-duty, each Addition to lower extremity, pelvic control, hip joint, | |
| | L2022 | | | adjustable flexion, each | |
| | L2624 | | | Addition to lower extremity, pelvic control, hip joint, | |
| | L2021 | | | adjustable flexion, extension, abduction control, each | |
| | L2627 | Y | Y | Addition to lower extremity, pelvic control, plastic, | |
| | | | | molded to patient model, reciprocating hip joint and | |
| | | | | cables | |
| | L2628 | Y | | Addition to lower extremity, pelvic control, metal | |
| | | | | frame, reciprocating hip joint and cables | |
| | L2630 | | | Addition to lower extremity, pelvic control, band and | |
| | L2640 | | | belt, unilateral Addition to lower extremity, pelvic control, band and | |
| | L2040 | | | belt, bilateral | |
| | L2650 | | | Addition to lower extremity, pelvic and thoracic | |
| | 2200 | | | control, gluteal pad, each | |
| | L2660 | | | Addition to lower extremity, thoracic control, thoracic | |
| | | | | band | |
| | | | | | |
| | L2670 | | | Addition to lower extremity, thoracic control, | |
| | 1.200 | | | paraspinal uprights | |
| | L2680 | | | Addition to lower extremity, thoracic control, lateral support uprights | |
| | L2750 | | Y | Addition to lower extremity orthosis, plating chrome | |
| | L2130 | | 1 | or nickel, per bar | |
| | L2755 | | Y | Addition to lower extremity orthosis, high strength, | |
| | | | | lightweight material, all hybrid lamination/prepreg | |
| | | | | composite, per segment | |

| Code | | | | | |
|----------------------|-------------------|----|-----|---|--|
| Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
| mulcators | L2760 | IA | Lic | Addition to lower extremity orthosis, extension, per | Comments |
| | L2700 | | | extension, per bar (for lineal adjustment for growth) | |
| | L2768 | Y | Y | Orthotic side bar disconnect device, per bar | |
| D | L2770 | _ | ¥ | Addition to lower extremity orthosis, any material, per | Removed |
| | | | | bar or joint | January 2010 |
| | L2780 | | Y | Addition to lower extremity orthosis, noncorrosive 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, kneecap, medial or lateral pull | |
| | L2810 | | | Addition to lower extremity orthosis, knee control, condylar pad | |
| | L2820 | | Y | Addition to lower extremity orthosis, soft interface for molded plastic, below knee section | |
| | L2830 | | Y | 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 | |
| # | L2860 | | | Addition to lower extremity joint, knee or ankle, concentric adjustable torsion style mechanism, each | |
| # | L2861 | | | Addition to lower extremity joint, knee or ankle, concentric adjustable torsion | Code added January 2010 |
| | L2999 | Y | Y | Lower extremity orthoses, not otherwise specified | <u>, </u> |
| | L3000 | Y | | Foot insert, removable, molded to patient model, "UCB" type, Berkeley Shell, each | See EPA criteria, pages E.5-E.7. |
| # | L3001 | | | Foot insert, removable, molded to patient model, Spenco, each. | |
| # | L3002 | | | Foot insert, removable, molded to patient model, Plastazote or equal, each | |
| # | L3003 | | | Foot insert, removable, molded to patient model, silicone gel, each | |
| # | L3010 | | | Foot insert, removable, molded to patient model, longitudinal arch support, each | |
| # | L3020 | | | Foot insert, removable, molded to patient model, longitudinal/metatarsal support, each | |
| | L3030 | Y | | Foot insert, removable, formed to patient foot, each | See EPA Criteria, pages E.5-E.7. |

| Code | D 1 | | | | D. II. / |
|----------------------|-------------------|----|-----|--|--|
| Status Indicators | Procedure Code | PA | Lic | Decorintion | Policy/ Comments |
| mulcators | L3031 | Y | Lic | Description Foot, insert/plate, removable, addition to lower | Comments |
| | L3031 | 1 | | extremity orthosis, high strength, lightweight material, | |
| | | | | all hybrid lamination/prepreg composite, each | |
| # | L3040 | | | Foot, arch support, removable, premolded, | |
| | | | | longitudinal, each | |
| # | L3050 | | | Foot, arch support, removable, premolded, metatarsal, each | |
| # | L3060 | | | Foot, arch support, removable, premolded longitudinal/metatarsal, each | |
| # | L3070 | | | Foot, arch support, nonremovable, attached to shoe, longitudinal, each | |
| # | L3080 | | | Foot, arch support, nonremovable, attached to shoe, metatarsal, each | |
| # | L3090 | | | Foot, arch support, nonremovable, attached to shoe, | |
| | | | | longitudinal/metatarsal, each | |
| | L3100 | | | Hallus-Valgus night dynamic splint | |
| | L3140 | | | Foot, abduction rotation bar, including shoes | |
| | L3150 | | | Foot, abduction rotation bar, without shoes | |
| # | L3160 | | | Foot, adjustable shoe-styled positioning device | |
| | L3170 | Y | | Foot, plastic, silicone or equal, heel stabilizer, each. | |
| # | L3201 | | | Orthopedic shoe, oxford with supinator or pronator, infant | |
| # | L3202 | | | Orthopedic shoe, oxford with supinator or pronator, child | |
| # | L3203 | | | Orthopedic shoe, oxford with supinator or pronator, junior | |
| # | L3204 | | | Orthopedic shoe, hightop with supinator or pronator, infant | |
| # | L3206 | | | Orthopedic shoe, hightop with supinator or pronator, child | |
| # | L3207 | | | Orthopedic shoe, hightop with supinator or pronator, junior | |
| # | L3208 | | | Surgical boot, each, infant | |
| # | L3209 | | | Surgical boot, each, child | |
| # | L3211 | | | Surgical boot, each, junior | |
| # | L3212 | | | Benesch boot, pair, infant | |
| # | L3213 | | | Benesch boot, pair, child | |
| # | L3214 | | | Benesch boot, pair, junior | |
| | L3215 | Y | | Orthopedic footwear, ladies shoe, oxford, each | See EPA criteria, pages E.5-E.7. |
| # | L3216 | | | Orthopedic footwear, ladies shoe, depth inlay, each | |
| # | L3217 | | | Orthopedic footwear, ladies shoe, hightop, depth inlay, each | |

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| Indicators | | PA | Lic | Description | Comments |
| | L3219 | Y | | Orthopedic footwear, mens shoe, oxford, each | See EPA |
| | | | | | criteria, pages E.5-E.7. |
| # | L3221 | | | Orthopedic footwear, mens shoe, each. depth inlay | |
| # | L3222 | | | Orthopedic footwear, mens shoe, hightop, depth inlay, each | |
| # | L3224 | | | Orthopedic footwear, woman's shoe, oxford, used as an integral part of brace (orthosis) | |
| # | L3225 | | | Orthopedic footwear, man's shoe, oxford, used as an integral part of a brace (orthosis) | |
| | L3230 | Y | | Orthopedic footwear, custom shoe, depth inlay, each. | |
| # | L3250 | | | Orthopedic footwear, custom molded shoe, removable inner mold, prosthetic shoe, each | |
| # | L3251 | | | Foot, shoe molded to patient model, silicone shoe, each | |
| # | L3252 | | | Foot, shoe molded to patient model, Plastazote (or similar), custom fabricated, each | |
| # | L3253 | | | Foot, molded shoe Plastazote (or similar), custom fitted, each | |
| # | L3254 | | | Nonstandard size or width | |
| # | L3255 | | | Nonstandard size or length | |
| # | L3257 | | | Orthopedic footwear, additional charge for split size | |
| # | L3260 | | | Surgical boot/shoe, each | |
| # | L3265 | | | Plastazote sandal, each | |
| # | L3300 | | | Lift, elevation, heel, tapered to metatarsals, per inch | |
| | L3310 | Y | | Lift, elevation, heel and sole, neoprene, per inch | See EPA criteria E.5-E.7. |
| | L3320 | Y | | Lift, elevation, heel and sole, cork, per inch | See EPA criteria E.5-E.7. |
| # | L3330 | | | Lift, elevation, metal extension (skate) | |
| # | L3332 | | | Lift, elevation, inside shoe, tapered, up to one-half inch | |
| | L3334 | Y | | Lift, elevation, heel, per inch | See EPA criteria E.5-E.7. |
| | L3340 | Y | | Heel wedge, SACH | _ |
| | L3350 | Y | | Heel wedge | |
| | L3360 | Y | | Sole wedge, outside sole | |
| # | L3370 | | | Sole wedge, between sole | |
| # | L3380 | | | Clubfoot wedge | |
| # | L3390 | | | Outflare wedge | |
| | L3400 | Y | | Metatarsal bar wedge, rocker | |
| | L3410 | Y | | Metatarsal bar wedge, between sole | |
| | L3420 | Y | | Full sole and heel wedge, between sole | |
| | L3430 | | | Heel, counter, plastic reinforced | |

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| Indicators | | PA | Lic | Description | Comments |
| # | L3440 | | | Heel, counter, leather reinforced | |
| # | L3450 | | | Heel, SACH cushion type | |
| # | L3455 | | | Heel, new leather, standard | |
| # | L3460 | | | Heel, new rubber, standard | |
| # | L3465 | | | Heel, Thomas with wedge | |
| # | L3470 | | | Heel, Thomas extended to ball | |
| # | L3480 | | | Heel, pad and depression for spur | |
| # | L3485 | | | Heel, pad, removable for spur | |
| # | L3500 | | | Orthopedic shoe addition, insole, leather | |
| # | L3510 | | | Orthopedic shoe addition, insole, rubber | |
| # | L3520 | | | Orthopedic shoe addition, insole, felt covered with | |
| | | | | leather | |
| # | L3530 | | | Orthopedic shoe addition, sole, half | |
| # | L3540 | | | Orthopedic shoe addition, sole, full | |
| # | L3550 | | | Orthopedic shoe addition, toe tap, standard | |
| # | L3560 | | | Orthopedic shoe addition, toe tap, horseshoe | |
| # | L3570 | | | Orthopedic shoe addition, special extension to instep | |
| | 23370 | | | (leather with eyelets) | |
| # | L3580 | | | Orthopedic shoe addition, convert instep to velcro | |
| | 23200 | | | closure | |
| # | L3590 | | | Orthopedic shoe addition, convert firm shoe counter to | |
| | | | | soft counter | |
| # | L3595 | | | Orthopedic shoe addition, March bar | |
| # | L3600 | | | Transfer of an orthosis from one shoe to another, | |
| | | | | caliper plate, existing | |
| # | L3610 | | | Transfer of an orthosis from one shoe to another, | |
| | | | | caliper plate, new | |
| | L3620 | | | Transfer of an orthosis from one shoe to another, solid | One in a 12- |
| | | | | stirrup, existing. | month period |
| | | | | | allowed without |
| | | | | | prior |
| | | | | | authorization |
| # | L3630 | | | Transfer of an orthosis from one shoe to another, solid | |
| | | | | stirrup, new | |
| # | L3640 | | | Transfer of an orthosis from one shoe to another, | |
| | | | | Dennis Browne splint (Riveton), both shoes | |
| # | L3649 | | | Orthopedic shoe, modification, addition or transfer, | |
| | | | | not otherwise specified | |
| | L3650 | | *** | SO, figure of eight design abduction restrainer, | |
| | | | | prefabricated, includes fitting and adjustment | |
| D | L3651 | - | *** | SO, single shoulder, elastic, prefabricated, includes | Removed |
| | | | | fitting and adjustment (e.g., neoprene, Lycra) | January 2010 |
| D | L3652 | - | *** | SO, double shoulder, elastic, prefabricated, includes | Removed |
| | | | | fitting and adjustment (e.g., neoprene, Lycra) | January 2010 |

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| Indicators | Code | PA | Lic | Description | Comments |
| | L3660 | | *** | SO, figure of eight design abduction restrainer, canvas and webbing, prefabricated, includes fitting and adjustment | |
| | L3670 | | *** | SO, acromio/clavicular (canvas and webbing type), prefabricated, includes fitting and adjustment | |
| | L3671 | Y | Y | SO, shoulder cap design, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| | L3672 | Y | Y | SO, abduction positioning (airplane design), thoracic component and support bar, without joints, may inleude soft interface, straps, custom | |
| | L3673 | Y | | SO, abduction positioning (airplane design), thoracic component and support bar, includes nontorsion joint/turnbuckle, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| # | L3675 | | | SO, vest type abduction restrainer, canvas webbing type, or equal, prefabricated, includes fitting and adjustment | |
| | L3677 | Y | Y | Shoulder orthosis, hard plastic, shoulder stabilizer, prefabricated, includes fitting and adjustment | |
| D | L3700 | - | *** | EO, elastic with stays, prefabricated, includes fitting and adjustment | Removed January 2010 |
| D | L3701 | - | <u>***</u> | EO, elastic, prefabricated, includes fitting and adjustment (e.g., neoprene, Lycra) | Removed January 2010 |
| | L3702 | Y | Y | EO, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| | L3710 | | *** | EO, elastic with metal joints, prefabricated, includes fitting and adjustment | |
| | L3720 | | | EO, double upright with forearm/arm cuffs, free motion, custom fabricated | |
| | L3730 | Y | Y | EO, double upright with forearm/arm cuffs, extension/flexion assist, custom fabricated | |
| | L3740 | Y | Y | EO, double upright with forearm/arm cuffs, adjustable position lock with active control, custom fabricated | |
| | L3760 | | | EO, with adjustable position, locking joints, prefabricated, includes fitting and adjustment, any type | |
| | L3762 | | *** | EO, rigid, without joints, includes soft interface material, prefabricated, includes fitting and adjustment | |
| | L3763 | Y | Y | EWHO, rigid, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| | L3764 | Y | Y | EWHO, includes one or more nontorsion joints, elastic bands, turnbuckles, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |

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| Indicators | Code | PA | Lic | Description | Comments |
| | L3765 | Y | Y | EWHFO, rigid, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| | L3766 | Y | Y | EWHFO, includes one or more nontorsion joints, elastic bands, turnbuckles, may include soft interface, straps, custom | |
| | L3806 | Y | Y | Wrist hand finger orthosis, includes one or more nontorsion joint(s), turnbuckles, elastic bands/springs, may include soft interface material, straps, custom fabricated, includes fitting and adjustment | |
| | L3807 | | | WHFO without joint(s), prefabricated, includes fitting and adjustment, any type | |
| | L3808 | Y | Y | Wrist hand finger orthosis, rigid without joints, may include soft interface material; straps, custom fabricated, includes fitting and adjustment | |
| # | L3891 | | | Addition to upper extremity joint, wrist or elbow, concentric adjustable | Code added January 2010 |
| | L3900 | | Y | WHFO, dynamic flexor hinge, reciprocal wrist extension/flexion, finger flexion/extension, wrist or finger driven, custom fabricated | · |
| | L3901 | Y | Y | WHFO, dynamic flexor hinge, reciprocal wrist extension/flexion, finger flexion/extension, cable driven, custom fabricated | |
| | L3904 | Y | Y | WHFO, external powered, electric, custom fabricated | |
| | L3905 | Y | Y | WHO, includes one or more nontorsion joints, elastic bands, turnbuckles, may include soft interface, straps, custom fabricated, includes | |
| | L3906 | | Y | WHO, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustment. | |
| | L3908 | | *** | WHO, wrist extension control cock-up, nonmolded, prefabricated, includes fitting and adjustments | |
| D | L3909 | - | <u>***</u> | Wrist orthosis, elastic, prefabricated, includes fitting and adjustment (e.g., neoprene, Lycra) | Removed January 2010 |
| D | L3911 | - | - | Wrist hand finger orthosis, elastic, prefabricated, includes fitting and adjustments (e.g., neoprene, Lycra) | Removed January 2010 |
| | L3912 | | *** | HFO, flexion glove with elastic finger control, prefabricated, includes fitting and adjustments | |
| | L3913 | Y | Y | HFO, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| | L3915 | Y | *** | Wrist hand orthosis, includes one or more nontorsion joint(s), elastic bands, turnbuckles, may include soft interface, straps, prefabricated, includes fitting and adjustment | |

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| | L3917 | | | HO, metacarpal fracture orthosis, prefabricated, includes fitting | COMMINA |
| | L3919 | Y | Y | Hand orthosis, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| | L3921 | Y | Y | Hand finger orthosis, includes one or more nontorsion joints, elastic bands, turnbuckles, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| | L3923 | | | HFO, without joints, may include soft interface, straps, prefabricated, includes fitting and adjustment | |
| | L3925 | Y | *** | Finger orthosis, proximal interphalangeal (Pip)/distal interphalangeal (dip), non torsion joint/spring, extension/flexion, may include soft interface material, prefabricated, includes fitting and adjustment. | |
| | L3927 | Y | | Finger orthosis, proximal interphalangeal (Pip)/distal interphalangeal (dip), without joint/spring, extension/flexion (e.g. static or ring type), may include soft interface material, prefabricated, includes fitting and adjustment. | |
| | L3929 | Y | | Hand finger orthosis, includes one or more nontorsion joint(s), turnbuckles, elastic bands/springs, may include soft interface material, straps, prefabricated, includes fitting and adjustment | |
| | L3931 | Y | | Wrist hand finger orthosis, includes one or more nontorsion joint(s), turnbuckles, elastic bands/springs, may include soft interface material, straps, prefabricated, includes fitting and adjustment | |
| | L3933 | Y | Y | Finger orthosis, without joints, may include soft interface, custom fabricated, includes fitting and adjustment | |
| | L3935 | Y | Y | Finger orthosis, nontorsion joint, may include soft interface, custom fabricated, includes fitting and adjustment | |
| | L3956 | Y | Y | Addition of joint to upper extremity orthosis, any material; per joint | |
| | L3960 | | | SEWHO, abduction positioning, airplane design, prefabricated, includes fitting and adjustments | |
| | L3961 | Y | Y | Shoulder elbow wrist hand orthosis, shoulder cap design, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| | L3962 | | | SEWHO, abduction positioning, Erb's palsey design, prefabricated, includes fitting and adjustments | |

| Code Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
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| # | L3964 | IA | Lic | SEO, mobile arm support attached to wheelchair, balanced, adjustable, prefabricated, includes fitting and adjustments | Comments |
| # | L3965 | | | SEO, mobile arm support attached to wheelchair, balanced, adjustable Rancho type, prefabricated, includes fitting and adjustments | |
| # | L3966 | | | SEO, mobile arm support attached to wheelchair, balanced, reclining, prefabricated, includes fitting and adjustments | |
| | L3967 | Y | | SEWHO, abduction positioning (airplane design), thoracic component and support bar, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustments | |
| # | L3968 | | | SEO, mobile arm support attached to wheelchair, balanced, friction arm support (friction dampening to proximal and distal joints), prefabricated, includes fitting and adjustments | |
| | L3969 | Y | | SEO, mobile arm support, monosuspension arm and hand support, overhead elbow forearm hand sling support, yoke type arm suspension support, prefabricated, includes fitting and adjustments | |
| | L3970 | | | SEO, addition to mobile arm support, elevating proximal arm | |
| | L3971 | Y | | SEWHO, shoulder cap design, includes one or more nontorsion joints, elastic bands, turnbuckles, may include soft interface | |
| | L3972 | | | SEO, addition to mobile arm support, offset or lateral rocker arm with elastic balance control | |
| | L3973 | Y | | SEWHO, abduction positioning (airplane design), thoracic component and support bar, includes one or more nontorsion joints | |
| | L3974 | | | SEO, addition to mobile arm support, supinator | |
| | L3975 | Y | Y | SEWHFO, shoulder cap design, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustment | |
| | L3976 | Y | | SEWHFO, abduction positioning (airplane design), thoracic component and support bar, without joints, may include soft interface, straps, custom fabricated, includes fitting and adjustments | |
| | L3977 | Y | | SEWHFO, shoulder cap design, includes one or more nontorsion joints, elastic bands, turnbuckles, may include soft interface, straps, custom fabricated, includes fitting and adjustments | |

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| Indicators | | PA | Lic | Description Company of the Company o | Comments |
| | L3978 | Y | | SEWHFO, abduction positioning (airplane design), thoracic component and support bar, includes one or | |
| | | | | more nontorsion | |
| | L3980 | | *** | Upper extremity fracture orthosis, humeral, | |
| | | | | prefabricated, includes fitting and adjustments | |
| | L3982 | | | Upper extremity fracture orthosis, radius/ulnar, | |
| | | | | prefabricated, includes fitting and adjustments | |
| | L3984 | | | Upper extremity fracture orthosis, wrist, prefabricated, includes fitting and adjustments | |
| | L3995 | | | Addition to upper extremity orthosis, sock, fracture or | |
| | | | | equal, each | |
| | L3999 | Y | Y | Upper limb orthosis, not otherwise specified | |
| | L4000 | Y | Y | Replace girdle for spinal orthosis (CTLSO or SO) | |
| | L4002 | Y | Y | Replacement strap, any orthosis, includes all | |
| | L4010 | | Y | components, any length, any type Replace trilateral socket brim | |
| | L4020 | | Y | Replace quadrilateral socket brim, molded to patient | |
| | L+020 | | 1 | model | |
| | L4030 | | Y | Replace quadrilateral socket brim, custom fitted | |
| | L4040 | | Y | Replace molded thigh lacer | |
| | L4045 | | Y | Replace nonmolded thigh lacer | |
| | L4050 | | Y | Replace molded calf lacer | |
| | L4055 | | Y | Replace nonmolded calf lacer | |
| | L4060 | | Y | Replace high roll cuff | |
| | L4070 | | Y | Replace proximal and distal upright for KAFO | |
| | L4080 | | Y | Replace metal bands KAFO, proximal thigh | |
| | L4090 | | Y | Replace metal bands KAFO-AFO, calf or distal thigh | |
| | L4100 | | Y | Replace leather cuff KAFO, proximal thigh | |
| | L4110 | | Y | Replace leather cuff KAFO–AFO, calf or distal thigh | |
| | L4130 | *7 | Y | Replace pretibial shell | |
| | L4205 | Y | Y | Repair of orthotic device, labor component, per 15 minutes | |
| | L4210 | Y | Y | Repair of orthotic device, repair or replace minor parts | |
| | L4350 | | *** | Pneumatic ankle control splint (e.g., aircast), prefabricated, includes fitting and adjustments | |
| | L4360 | Y | | Pneumatic ankle foot orthosis, with or without joints, prefabricated, includes fitting and adjustments | |
| | L4370 | Y | *** | Pneumatic full leg splint (e.g., aircast), prefabricated, includes fitting and adjustments | |
| | L4380 | | *** | Pneumatic knee splint (e.g., aircast), prefabricated, includes fitting and adjustments | |
| | L4386 | Y | *** | Non-pneumatic walking splint, with or without joints, prefabricated, includes fitting and adjustments | |
| # | L4392 | | | Replacement soft interface material, static AFO | |

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| Indicators # | Code L4394 | PA | Lic | Description Replace soft interface material, foot drop splint | Comments |
| π | L4396 | Y | | Static ankle foot orthosis, including soft interface material, adjustable for fit, for positioning, pressure reduction, may be used for minimal ambulation, prefabricated, includes fitting and adjustments | |
| # | L4398 | | | Foot drop splint, recumbent positioning device, prefabricated, includes fitting and adjustments | |
| | L5000 | | Y | Partial foot, shoe insert with longitudinal arch, toe filler | |
| | L5010 | | Y | Partial foot, molded socket, ankle height, with toe filler | |
| | L5020 | | Y | Partial foot, molded socket, tibial tubercle height, with toe filler | |
| | L5050 | | Y | Ankle, Symes, molded socket, SACH Foot | |
| | L5060 | Y | Y | Ankle, Symes, metal frame, molded leather socket, articulated ankle/foot | |
| | L5100 | | Y | Below knee, molded socket, shin, SACH foot | |
| | L5105 | Y | Y | Below knee, plastic socket, joints and thigh lacer, SACH foot | |
| | L5150 | Y | Y | Knee disarticulation (or through knee), molded socket, external knee joints, shin, SACH foot | |
| | L5160 | Y | Y | Knee disarticulation (or through knee), molded socket, bent knee configuration, external knee joints, shin, SACH foot | |
| | L5200 | | Y | Above knee, molded socket, single axis constant friction knee, shin, SACH foot | |
| | L5210 | | Y | Above knee, short prosthesis, no knee joint ("stubbies"), with foot blocks, no ankle joints, each | |
| | L5220 | Y | Y | Above knee, short prosthesis, no knee joint ("stubbies"), with articulated ankle/foot, dynamically aligned, each | |
| | L5230 | Y | Y | Above knee, for proximal femoral focal deficiency, constant friction knee, shin, SACH foot | |
| | L5250 | Y | Y | Hip disarticulation, Canadian type; molded socket, hip joint, single axis constant friction knee, shin, SACH foot | |
| | L5270 | Y | Y | Hip disarticulation, tilt table type; molded socket, locking hip joint, single axis constant friction knee, shin, SACH foot | |
| | L5280 | Y | Y | Hemipelvectomy, Canadian type; molded socket, hip joint, single axis constant friction knee, shin, SACH foot | |
| | L5301 | | Y | Below knee, molded socket, shin, SACH foot, endoskeletal system | |

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| 220200020 | L5311 | | Y | Knee disarticulation (or through knee), molded socket, external knee joints, shin, SACH foot, endoskeletal system | |
| | L5321 | | Y | Above knee, molded socket, open end, SACH foot, endoskeletal system, single axis knee | |
| | L5331 | | Y | Hip disarticulation, Canadian type, molded socket, endoskeletal system, hip joint, single axis knee, SACH foot | |
| | L5341 | | Y | Hemipelvectomy, Canadian type, molded socket, endoskeletal system, hip joint, single axis knee, SACH foot | |
| | L5400 | | Y | Immediate postsurgical or early fitting, application of initial rigid dressing, including fitting, alignment, suspension, and one cast change, below knee | |
| | L5410 | | Y | Immediate postsurgical or early fitting, application of initial rigid dressing, including fitting, alignment and suspension, below knee, each additional cast change and realignment | |
| | L5420 | | Y | Immediate postsurgical or early fitting, application of initial rigid dressing, including fitting, alignment and suspension and one cast change AK or knee disarticulation | |
| | L5430 | | Y | Immediate postsurgical or early fitting, application of initial rigid dressing, including fitting, alignment and suspension, AK or knee disarticulation, each additional cast change and realignment | |
| | L5450 | | Y | Immediate postsurgical or early fitting, application of nonweight bearing rigid dressing, below knee | |
| | L5460 | | Y | Immediate postsurgical or early fitting, application of nonweight bearing rigid dressing, above knee | |
| | L5500 | | Y | Initial, below knee PTB type socket, non-alignable system, pylon, no cover, SACH foot, plaster socket, direct formed | |
| | L5505 | Y | Y | Initial, above knee – knee disarticulation, ischial level socket, non-alignable system, pylon, no cover, SACH foot plaster socket, direct formed | |
| | L5510 | | Y | Preparatory, below knee PTB type socket, non- alignable system, pylon, no cover, SACH foot, plaster socket, molded to model | Limit one per client per lifetime per limb |
| | L5520 | | Y | Preparatory, below knee PTB type socket, non- alignable system, pylon, no cover, SACH foot, thermoplastic or equal, direct formed | Limit one per client per lifetime per limb |

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| Indicators | | PA | Lic | Description | Comments |
| | L5530 | Y | Y | Preparatory, below knee PTB type socket, non- | |
| | | | | alignable system, pylon, no cover, SACH foot, | |
| | | | | thermoplastic or equal, molded to model | |
| | L5535 | Y | Y | Preparatory, below knee PTB type socket, non- | |
| | | | | alignable system, pylon, no cover, SACH foot, | |
| | T 5540 | *7 | X 7 | prefabricated, adjustable open end socket | |
| | L5540 | Y | Y | Preparatory, below knee PTB type socket, non- | |
| | | | | alignable system, pylon, no cover, SACH foot, | |
| | 1.55(0) | Y | Y | laminated socket, molded to model | |
| | L5560 | ĭ | ĭ | Preparatory, above knee – knee disarticulation, ischial | |
| | | | | level socket, non-alignable system, pylon, no cover, SACH foot, plaster socket, molded to model | |
| | L5570 | Y | Y | Preparatory, above knee - knee disarticulation, ischial | |
| | L3370 | 1 | 1 | level socket, non-alignable system, pylon, no cover, | |
| | | | | SACH foot, thermoplastic or equal, direct formed | |
| | L5580 | Y | Y | Preparatory, above knee – knee disarticulation, ischial | |
| | L3300 | 1 | • | level socket, non-alignable system, pylon, no cover, | |
| | | | | SACH foot, thermoplastic or equal, molded to model | |
| | L5585 | Y | Y | Preparatory, above knee – knee disarticulation, ischial | |
| | | | | level socket, non-alignable system, pylon, no cover, | |
| | | | | SACH foot, prefabricated adjustable open end socket | |
| | L5590 | Y | Y | Preparatory, above knee – knee disarticulation, ischial | |
| | | | | level socket, non-alignable system, pylon, no cover, | |
| | | | | SACH foot, laminated socket, molded to model | |
| | L5595 | Y | Y | Preparatory, hip disarticulation – hemipelvectomy, | |
| | | | | pylon, no cover, SACH foot, thermoplastic or equal, | |
| | | | | molded to patient model | |
| | L5600 | Y | Y | Preparatory, hip disarticulation – hemipelvectomy, | |
| | | | | pylon, no cover, SACH foot, laminated socket, | |
| | | | | molded to patient model | |
| | L5610 | Y | Y | Addition to lower extremity, endoskeletal system, | |
| | | | | above knee, hydracadence system | |
| | L5611 | Y | Y | Addition to lower extremity, endoskeletal system, | |
| | | | | above knee - knee disarticulation, 4-bar linkage, with | |
| | I 5(12 | 3.7 | 37 | friction swing phase control | |
| | L5613 | Y | Y | Addition to lower extremity, endoskeletal system, | |
| | | | | above knee - knee disarticulation, 4-bar linkage, with | |
| | L5614 | Y | Y | hydraulic swing phase control Addition to lower extremity, endoskeletal system, | |
| | L3014 | 1 | 1 | above knee - knee disarticulation, 4-bar linkage, with | |
| | | | | pneumatic swing phase control | |
| | L5616 | | Y | Addition to lower extremity, endoskeletal system, | |
| | L3010 | | 1 | above knee, universal multiplex system, friction swing | |
| | | | | phase control | |
| | L5617 | | Y | † * | |
| | | | • | | |
| | L5617 | | Y | Addition to lower extremity, quick change self- aligning unit, above or below knee, each | |

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| Indicators | Code | PA | Lic | Description | Comments |
| | L5618 | | Y | Addition to lower extremity, test socket, Symes | |
| | L5620 | | Y | Addition to lower extremity, test socket, below knee | |
| | L5622 | | Y | Addition to lower extremity, test socket, knee | |
| | 1.5604 | | X 7 | disarticulation | |
| | L5624 | | Y | Addition to lower extremity, test socket, above knee | |
| | L5626 | | Y | Addition to lower extremity, test socket, hip disarticulation | |
| | L5628 | | Y | Addition to lower extremity, test socket, hemipelvectomy | |
| | L5629 | | Y | Addition to lower extremity, below knee, acrylic socket | |
| | L5630 | | Y | Addition to lower extremity, Symes type, expandable wall socket | |
| | L5631 | | Y | Addition to lower extremity, above knee or knee disarticulation, acrylic socket | |
| | L5632 | | Y | Addition to lower extremity, Symes type, PTB brim design socket | |
| | L5634 | | Y | Addition to lower extremity, Symes type, posterior opening (Canadian) socket | |
| | L5636 | | Y | Addition to lower extremity, Symes type, medial opening socket | |
| | L5637 | | Y | Addition to lower extremity, below knee, total contact | |
| | L5638 | Y | Y | Addition to lower extremity, below knee, leather | |
| | | | | socket | |
| | L5639 | Y | Y | Addition to lower extremity, below knee, wood socket | |
| | L5640 | Y | Y | Addition to lower extremity, knee disarticulation, leather socket | |
| | L5642 | Y | Y | Addition to lower extremity, above knee, leather socket | |
| | L5643 | Y | Y | Addition to lower extremity, hip disarticulation, flexible inner socket, external frame | |
| | L5644 | Y | Y | Addition to lower extremity, above knee, wood socket | |
| | L5645 | Y | Y | Addition to lower extremity, below knee, flexible inner socket, external frame | |
| | L5646 | Y | Y | Addition to lower extremity, below knee, air cushion | |
| | L5647 | Y | Y | Addition to lower extremity, below knee, suction | |
| | L5648 | Y | Y | Socket Addition to lower extremity, above knee, air cushion | |
| | L5649 | | Y | Socket Addition to lower extremity, ischial | |
| | | | | containment/narrow M-L socket | |
| | L5650 | | Y | Addition to lower extremity, total contact, above knee or knee disarticulation socket | |
| | L5651 | | Y | Addition to lower extremity, above knee, flexible | |

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| | 0000 | | | inner socket, external frame | 00111110110 |
| | L5652 | | Y | Addition to lower extremity, suction suspension, above knee or knee disarticulation socket | |
| | L5653 | | Y | Addition to lower extremity, knee disarticulation, expandable wall socket | |
| | L5654 | | Y | Addition to lower extremity, socket insert, Symes (Kemblo, Pelite, Aliplast, Plastazote or equal) | |
| | L5655 | | Y | Addition to lower extremity, socket insert, below knee (Kemblo, Pelite, Aliplast, Plastazote or equal) | |
| | L5656 | | Y | Addition to lower extremity, socket insert, knee disarticulation (Kemblo, Pelite, Aliplast, Plastazote or equal) | |
| | L5658 | Y | Y | Addition to lower extremity, socket insert, above knee (Kemblo, Pelite, Aliplast, Plastazote or equal) | |
| | L5661 | Y | Y | Addition to lower extremity, socket insert, multidurometer, Symes | |
| | L5665 | | Y | Addition to lower extremity, socket insert, multidurometer, below knee | |
| | L5666 | | Y | Addition to lower extremity, below knee, cuff suspension | |
| | L5668 | | Y | Addition to lower extremity, below knee, molded distal cushion | |
| | L5670 | | Y | Addition to lower extremity, below knee, molded supracondylar suspension (PTS or similar) | |
| | L5671 | | Y | Addition to lower extremity, below knee/above knee suspension locking mechanism (shuttle, lanyard or equal), excludes socket insert | |
| | L5672 | | Y | Addition to lower extremity, below knee, removable medial brim suspension | |
| | L5673 | | Y | Addition to lower extremity, below knee/above knee, custom fabricated from existing mold or prefabricated, socket insert, silicone gel, elastomeric or equal, for use with locking mechanism | |
| | L5676 | | Y | Addition to lower extremity, below knee, knee joints, single axis, pair | |
| | L5677 | Y | Y | Addition to lower extremity, below knee, knee joints, polycentric, pair | |
| | L5678 | | Y | Addition to lower extremity, below knee, joint covers, pair | |
| | L5679 | | Y | Addition to lower extremity, below knee/above knee, custom fabricated from existing mold or prefabricated, socket insert, silicone gel, elastomeric or equal, not for use with locking mechanism | |
| | L5680 | | Y | Addition to lower extremity, below knee, thigh lacer, nonmolded | |

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| Indicators | Code | PA | Lic | Description | Comments |
| | L5681 | Y | Y | Addition to lower extremity, below knee/above knee, custom fabricated socket insert for congenital or atypical traumatic amputee, silicone gel, elastomeric or equal, for use with or without locking mechanism, initial only (for other than initial, use code L5673 or L5679) | |
| | L5682 | Y | Y | Addition to lower extremity, below knee, thigh lacer, gluteal/ischial, molded | |
| | L5683 | Y | Y | Addition to lower extremity, below knee/above knee, custom fabricated socket insert for other than congenital or atypical traumatic amputee, silicone gel, elastomeric or equal, for use with or without locking mechanism, initial only (for other than initial, use code L5673 or L5679) | |
| | L5684 | | Y | Addition to lower extremity, below knee, fork strap | |
| | L5685 | | Y | Addition to lower extremity prosthesis, below knee, suspension/sealing sleeve, with or without valve, any material, each | |
| | L5686 | | Y | Addition to lower extremity, below knee, back check (extension control) | |
| | L5688 | | Y | Addition to lower extremity, below knee, waist belt, webbing | |
| | L5690 | | Y | Addition to lower extremity, below knee, waist belt, padded and lined | |
| | L5692 | | Y | Addition to lower extremity, above knee, pelvic control belt, light | |
| | L5694 | | Y | Addition to lower extremity, above knee, pelvic control belt, padded and lined | |
| | L5695 | | Y | Addition to lower extremity, above knee, pelvic control, sleeve suspension, neoprene or equal, each | |
| | L5696 | | Y | Addition to lower extremity, above knee or knee disarticulation, pelvic joint | |
| | L5697 | | Y | Addition to lower extremity, above knee or knee disarticulation, pelvic band | |
| | L5698 | | Y | Addition to lower extremity, above knee or knee disarticulation, Silesian bandage | |
| | L5699 | | Y | All lower extremity prostheses, shoulder harness | |
| | L5700 | | Y | Replacement, socket, below knee, molded to patient model | Limit one per client per year |
| | L5701 | | Y | Replacement, socket, above knee/knee disarticulation, including attachment plate, molded to patient model | Limit one per client per year |
| | L5702 | Y | Y | Replacement, socket, hip disarticulation, including hip joint, molded to patient model | • • |
| | L5703 | Y | Y | Ankle, symes, molded to patient model, socket without solid ankle cushion heel (sach) foot, replacement only | |

| | Procedure | | | | Policy/ |
|------------|-----------|----|-----|--|----------|
| Indicators | | PA | Lic | Description | Comments |
| | L5704 | Y | Y | Custom shaped protective cover, below knee | |
| | L5705 | Y | Y | Custom shaped protective cover, above knee | |
| | L5706 | Y | Y | Custom shaped protective cover, knee disarticulation | |
| | L5707 | Y | Y | Custom shaped protective cover, hip disarticulation | |
| | L5710 | | Y | Addition, exoskeletal knee-shin system, single axis, manual lock | |
| | L5711 | | Y | Addition, exoskeletal knee-shin system, single axis, manual lock, ultra-light material | |
| | L5712 | | Y | Addition, exoskeletal knee-shin system, single axis, friction swing and stance phase control (safety knee) | |
| | L5714 | | Y | Addition, exoskeletal knee-shin system, single axis, variable friction swing phase control | |
| | L5716 | Y | Y | Addition, exoskeletal knee-shin system, polycentric, mechanical stance phase lock | |
| | L5718 | Y | Y | Addition, exoskeletal knee-shin system, polycentric, friction swing and stance phase control | |
| | L5722 | | Y | Addition, exoskeletal knee-shin system, single axis, pneumatic swing, friction stance phase control | |
| | L5724 | Y | Y | Addition, exoskeletal knee-shin system, single axis, fluid swing phase control | |
| | L5726 | Y | Y | Addition, exoskeletal knee-shin system, single axis, external joints, fluid swing phase control | |
| | L5728 | Y | Y | Addition, exoskeletal knee-shin system, single axis, fluid swing and stance phase control | |
| | L5780 | | Y | Addition, exoskeletal knee-shin system, single axis, pneumatic/hydra pneumatic swing phase control | |
| | L5781 | Y | Y | Addition to lower limb prosthesis, vacuum pump, residual limb volume management and moisture evacuation system | |
| | L5782 | Y | Y | Addition to lower limb prosthesis, vacuum pump, residual limb volume management and moisture evacuation system, heavy duty | |
| | L5785 | | Y | Addition, exoskeletal system, below knee, ultra-light material (titanium, carbon fiber or equal) | |
| | L5790 | | Y | Addition, exoskeletal system, above knee, ultra-light material (titanium, carbon fiber or equal) | |
| | L5795 | | Y | Addition, exoskeletal system, hip disarticulation, ultra-light material (titanium carbon fiber or equal) | |
| | L5810 | | Y | Addition, endoskeletal knee-shin system, single axis, manual lock | |
| | L5811 | | Y | Addition, endoskeletal knee-shin system, single axis, manual lock, ultra-light material | |
| | L5812 | | Y | Addition, endoskeletal knee-shin system, single axis, friction swing and stance phase control (safety knee) | |

| Code | | | | | D. W. / |
|----------------------|-------------------|-----|-----|---|---------------------|
| Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
| mulcators | L5814 | Y | Y | Addition, endoskeletal knee-shin system, polycentric, | Comments |
| | | | | hydraulic swing phase control, mechanical stance | |
| | | | | phase lock | |
| | L5816 | | Y | Addition, endoskeletal knee-shin system, polycentric, | |
| | | | | mechanical stance phase lock | |
| | L5818 | | Y | Addition, endoskeletal knee-shin system, polycentric, friction swing and stance phase control | |
| | L5822 | | Y | Addition, endoskeletal knee-shin system, single axis, pneumatic swing, friction stance phase control | |
| | L5824 | | Y | Addition, endoskeletal knee-shin system, single axis, fluid swing phase control | |
| | L5826 | Y | Y | Addition, endoskeletal knee-shin system, single axis, | |
| | | | | hydraulic swing phase control, with miniature high | |
| | | | | activity frame | |
| | L5828 | Y | Y | Addition, endoskeletal knee-shin system, single axis, | |
| | ¥ 5020 | * 7 | * 7 | fluid swing and stance phase control | |
| | L5830 | Y | Y | Addition, endoskeletal knee-shin system, single axis, pneumatic/swing phase control | |
| | L5840 | Y | Y | Addition, endoskeletal knee-shin system, 4-bar | |
| | 23010 | - | • | linkage or multiaxial, pneumatic swing phase control | |
| # | L5845 | | | Addition, endoskeletal knee-shin system, stance | |
| | | | | flexion feature, adjustable | |
| | L5848 | Y | Y | Addition to, endoskeletal, knee-shin system, hydraulic | |
| | | | | stance extension, dampening feature, adjustable | |
| | L5850 | | Y | Addition, endoskeletal system, above knee or hip | |
| | L5855 | | Y | disarticulation, knee extension assist Addition, endoskeletal system, hip disarticulation, | |
| | L3033 | | 1 | mechanical hip extension assist | |
| # | L5856 | Y | Y | Addition to lower extremity prosthesis, endoskeletal | |
| | | | | knee-shin system, microprocessor control feature, | |
| | | | | swing and stance phase, includes electronic sensor(s), | |
| | | | | any type | |
| | L5857 | Y | Y | Addition to lower extremity prosthesis, endoskeletal | |
| | | | | knee-shin system, microprocessor control feature, swing phase only, includes electronic sensor(s), any | |
| | | | | type | |
| # | L5858 | | | Addition to lower extremity prosthesis, endoskeletal | |
| | | | | knee shin system, microprocessor control feature, | |
| | | | | stance phase only, includes electronic sensor(s), any | |
| | Y 5010 | | ** | type | |
| | L5910 | | Y | Addition, endoskeletal system, below knee, alignable | |
| | L5920 | | Y | System Addition, endoskeletal system, above knee or hip | |
| | L3720 | | 1 | disarticulation, alignable system | |
| | L5925 | | Y | Addition, endoskeletal system, above knee, knee | |
| | | | | disarticulation or hip disarticulation, manual lock | |

| Code | | | | | |
|------------|-----------|----|------------|---|---------------|
| | Procedure | | - . | 5 | Policy/ |
| Indicators | | PA | Lic | Description | Comments |
| # | L5930 | | | Addition, endoskeletal system, high activity knee control frame | |
| | L5940 | Y | Y | Addition, endoskeletal system, below knee, ultra-light | |
| | L3940 | 1 | 1 | material (titanium, carbon fiber or equal) | |
| | L5950 | Y | Y | Addition, endoskeletal system, above knee, ultra-light | |
| | L3730 | 1 | 1 | material (titanium, carbon fiber or equal) | |
| | L5960 | Y | Y | Addition, endoskeletal system, hip disarticulation, | |
| | 20,00 | _ | - | ultra-light material (titanium, carbon fiber or equal) | |
| | L5962 | Y | Y | Addition, endoskeletal system, below knee, flexible | |
| | | | | protective outer surface covering system | |
| | L5964 | Y | Y | Addition, endoskeletal system, above knee, flexible | |
| | | | | protective outer surface covering system | |
| | L5966 | Y | Y | Addition, endoskeletal system, hip disarticulation, | |
| | | | | flexible protective outer surface covering system | |
| | L5968 | Y | Y | Addition to lower limb prosthesis, multiaxial ankle | |
| | | | | with swing phase action dorsiflexion feature | |
| | L5970 | Y | Y | All lower extremity prostheses, foot, external keel, | |
| | | | | SACH foot | |
| | L5971 | Y | Y | All lower extremity prosthesis, solid ankle cushion | |
| | 1.5070 | | 3 7 | heel (sach) foot, replacement only | |
| | L5972 | | Y | All lower extremity prostheses, flexible keel foot | |
| # | L5973 | | | (safe, sten, bock dynamic or equal) Endoskeletal ankle foot system, microprocessor | Code added |
| # | L3913 | | | controlled feature, dorsiflexion | January 2010 |
| | L5974 | | Y | All lower extremity prostheses, foot, single axis | Juliuary 2010 |
| | 20,7. | | - | ankle/foot | |
| | L5975 | | Y | All lower extremity prosthesis, combination single | |
| | | | | axis and flexible keel foot | |
| | L5976 | | Y | All lower extremity prostheses, energy storing foot | |
| | | | | (Seattle carbon copy II or equal) | |
| | L5978 | | Y | All lower extremity prostheses, foot, multi-axial | |
| | | | | ankle/foot | |
| | L5979 | Y | Y | All lower extremity prostheses, multi-axial ankle, | |
| | T 5000 | ** | X 7 | dynamic response foot, one piece system | |
| | L5980 | Y | Y | All lower extremity prostheses, flex-foot system | |
| | L5981 | Y | Y | All lower extremity prostheses, flex-walk system or | |
| | 1.5092 | Y | Y | equal All exoskeletal lower extremity prostheses, axial | |
| | L5982 | 1 | 1 | rotation unit | |
| | L5984 | Y | Y | All endoskeletal lower extremity prostheses, axial | |
| | L3707 | 1 | 1 | rotation unit | |
| | L5985 | Y | Y | All endoskeletal lower extremity prostheses, dynamic | |
| | | | - | prosthetic pylon | |
| | L5986 | Y | Y | All lower extremity prostheses, multi-axial rotation | |
| | | | | unit (MCP or equal) | |

| Code Status | Procedure | | | | Policy/ |
|----------------|-----------|----|-----|---|----------|
| Indicators | | PA | Lic | Description | Comments |
| # | L5987 | | | All lower extremity prostheses, shank foot system with vertical loading pylon | |
| | L5988 | Y | Y | Addition to lower limb prosthesis, vertical shock reducing pylon feature | |
| | L5990 | Y | Y | Addition to lower extremity prosthesis, user adjustable heel height | |
| | L5993 | Y | Y | Addition to lower extremity prosthesis, heavy duty feature, foot only, (for patient weight greater than 300 lbs) Discontinued 1/1/09 | |
| | L5994 | Y | Y | Addition to lower extremity prosthesis, heavy duty feature, knee only, (for patient weight greater than 300 lbs) Discontinued 1/1/09 | |
| | L5995 | Y | Y | Addition to lower extremity prosthesis, heavy duty feature (for patient weight greater than 300 lbs) Discontinued 1/1/09 | |
| | L5999 | Y | Y | Lower extremity prosthesis, not otherwise specified | |
| | L6000 | Y | Y | Partial hand, Robin-Aids, thumb remaining (or equal) | |
| | L6010 | Y | Y | Partial hand, Robin-Aids, little and/or ring finger remaining (or equal) | |
| | L6020 | Y | Y | Partial hand, Robin-Aids, no finger remaining (or equal) | |
| | L6025 | Y | Y | Transcarpal/metacarpal or partial hand disarticulation prosthesis, external power, self-suspended, inner socket with removable forearm section, electrodes and cables, two batteries, charger, myoelectric control of terminal device | |
| | L6050 | | Y | Wrist disarticulation, molded socket, flexible elbow hinges, triceps pad | |
| | L6055 | Y | Y | Wrist disarticulation, molded socket with expandable interface, flexible elbow hinges, triceps pad | |
| | L6100 | | Y | Below elbow, molded socket, flexible elbow hinge, triceps pad | |
| | L6110 | | Y | Below elbow, molded socket (Muenster or Northwestern suspension types) | |
| | L6120 | Y | Y | Below elbow, molded double wall split socket, step-up hinges, half cuff | |
| | L6130 | Y | Y | Below elbow, molded double wall split socket, stump activated locking hinge, half cuff | |
| | L6200 | | Y | Elbow disarticulation, molded socket, outside locking hinge, forearm | |
| | L6205 | Y | Y | Elbow disarticulation, molded socket with expandable interface, outside locking hinges, forearm | |
| | L6250 | | Y | Above elbow, molded double wall socket, internal locking elbow, forearm | |
| | L6300 | | Y | Shoulder disarticulation, molded socket, shoulder | |

| | Procedure | | | | Policy/ |
|------------|-----------|----|-----|---|----------|
| Indicators | Code | PA | Lic | Description | Comments |
| | | | | bulkhead, humeral section, internal locking elbow, forearm | |
| | L6310 | Y | Y | Shoulder disarticulation, passive restoration (complete prosthesis) | |
| | L6320 | Y | Y | Shoulder disarticulation, passive restoration (shoulder cap only) | |
| | L6350 | Y | Y | Interscapular thoracic, molded socket, shoulder bulkhead, humeral section, internal locking elbow, forearm | |
| | L6360 | Y | Y | Interscapular thoracic, passive restoration (complete prosthesis) | |
| | L6370 | Y | Y | Interscapular thoracic, passive restoration (shoulder cap only) | |
| | L6380 | | Y | Immediate postsurgical or early fitting, application of initial rigid dressing, including fitting alignment and suspension of components, and one cast change, wrist disarticulation or below elbow | |
| | L6382 | | Y | Immediate postsurgical or early fitting, application of initial rigid dressing including fitting alignment and suspension of components, and one cast change, elbow disarticulation or above elbow | |
| | L6384 | | Y | Immediate postsurgical or early fitting, application of initial rigid dressing including fitting alignment and suspension of components, and one cast change, shoulder disarticulation or interscapular thoracic | |
| | L6386 | | Y | Immediate postsurgical or early fitting, each additional cast change and realignment | |
| | L6388 | | Y | Immediate postsurgical or early fitting, application of rigid dressing only | |
| | L6400 | | Y | Below elbow, molded socket, endoskeletal system, including soft prosthetic tissue shaping | |
| | L6450 | Y | Y | Elbow disarticulation, molded socket, endoskeletal system, including soft prosthetic tissue shaping | |
| | L6500 | | Y | Above elbow, molded socket, endoskeletal system, including soft prosthetic tissue shaping | |
| | L6550 | | Y | Shoulder disarticulation, molded socket, endoskeletal system, including soft prosthetic tissue shaping | |
| | L6570 | | Y | Interscapular thoracic, molded socket, endoskeletal system, including soft prosthetic tissue shaping | |
| | L6580 | Y | Y | Preparatory, wrist disarticulation or below elbow, single wall plastic socket, friction wrist, flexible elbow hinges, figure of eight harness, humeral cuff, Bowden cable control, USMC or equal pylon, no cover, molded to patient model | |
| | L6582 | Y | Y | Preparatory, wrist disarticulation or below elbow, single wall socket, friction wrist, flexible elbow | |

| Code Status | Procedure | | | | Policy/ |
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| Indicators | Code | PA | Lic | Description | Comments |
| | | | | hinges, figure of eight harness, humeral cuff, Bowden cable control, USMC or equal pylon, no cover, direct formed | |
| | L6584 | Y | Y | Preparatory, elbow disarticulation or above elbow, single wall plastic socket, friction wrist, locking elbow, figure of eight harness, fair lead cable control, USMC or equal pylon, no cover, molded to patient model | |
| | L6586 | Y | Y | Preparatory, elbow disarticulation or above elbow, single wall socket, friction wrist, locking elbow, figure of eight harness, fair lead cable control, USMC or equal pylon, no cover, direct formed | |
| | L6588 | Y | Y | Preparatory, shoulder disarticulation or interscapular thoracic, single wall plastic socket, shoulder joint, locking elbow, friction wrist, chest strap, fair lead cable control, USMC or equal pylon, no cover, molded to patient model | |
| | L6590 | Y | Y | Preparatory, shoulder disarticulation or interscapular thoracic, single wall socket, shoulder joint, locking elbow, friction wrist, chest strap, fair lead cable control, USMC or equal pylon, no cover, direct formed | |
| | L6600 | | Y | Upper extremity additions, polycentric hinge, pair | |
| | L6605 | | Y | Upper extremity additions, single pivot hinge, pair | |
| | L6610 | | Y | Upper extremity additions, flexible metal hinge, pair | |
| | L6611 | Y | Y | Addition to upper extremity prosthesis, external powered, additional switch, any type | |
| | L6615 | | Y | Upper extremity addition, disconnect locking wrist unit | |
| | L6616 | | Y | Upper extremity addition, additional disconnect insert for locking wrist unit, each | |
| | L6620 | | Y | Upper extremity addition, flexion-friction wrist unit | |
| | L6621 | Y | Y | Upper extremity prosthesis addition, flexion/extension wrist with or without friction, for use with external powered terminal device | |
| | L6623 | Y | Y | Upper extremity addition, spring assisted rotational wrist unit with latch release | |
| | L6624 | Y | Y | Upper extremity addition, flexion/extension and rotation wrist unit | |
| | L6625 | Y | Y | Upper extremity addition, rotational wrist unit with cable lock | |
| | L6628 | | Y | Upper extremity addition, quick disconnect hook adapter, Otto Bock or equal | |
| | L6629 | | Y | Upper extremity addition, quick disconnect lamination collar with coupling piece, Otto Bock or equal | |
| | L6630 | | Y | Upper extremity addition, stainless steel, any wrist | |

| | Procedure | | | | Policy/ |
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| Indicators | | PA | Lic | Description | Comments |
| | L6632 | | Y | Upper extremity addition, latex suspension sleeve, each | |
| | L6635 | | Y | Upper extremity addition, lift assist for elbow | |
| | L6637 | Y | Y | Upper extremity addition, nudge control elbow lock | |
| | L6638 | Y | Y | Upper extremity addition to prosthesis, electric locking feature, only for use with manually powered elbow | |
| D | L6639 | ¥ | ¥ | Upper extremity addition, heavy duty feature, any elbow | Removed January 2010 |
| | L6640 | Y | Y | Upper extremity additions, shoulder abduction joint, pair | |
| | L6641 | Y | Y | Upper extremity addition, excursion amplifier, pulley type | |
| | L6642 | Y | Y | Upper extremity addition, excursion amplifier, lever type | |
| | L6645 | | Y | Upper extremity addition, shoulder flexion-abduction joint, each | |
| | L6646 | Y | Y | Upper extremity addition, shoulder joint, multipositional locking, flexion, adjustable abduction friction control, for use with body powered or external powered system | |
| | L6647 | | Y | Upper extremity addition, shoulder lock mechanism, body powered actuator | |
| | L6648 | Y | Y | Upper extremity addition, shoulder lock mechanism, external powered actuator | |
| | L6650 | | Y | Upper extremity addition, shoulder universal joint, each | |
| | L6655 | | Y | Upper extremity addition, standard control cable, extra | |
| | L6660 | | Y | Upper extremity addition, heavy duty control cable | |
| | L6665 | | Y | Upper extremity addition, Teflon, or equal, cable lining | |
| | L6670 | | Y | Upper extremity addition, hook to hand, cable adapter | |
| | L6672 | | Y | Upper extremity addition, harness, chest or shoulder, saddle type | |
| | L6675 | | Y | Upper extremity addition, harness, figure of eight type, for single control | |
| | L6676 | | Y | Upper extremity addition, harness, figure of eight type, for dual control | |
| | L6677 | Y | Y | Upper extremity addition, harness, triple control, simultaneous operation of terminal device and elbow. | |
| | L6680 | | Y | Upper extremity addition, test socket, wrist disarticulation or below elbow | |
| | L6682 | | Y | Upper extremity addition, test socket, elbow disarticulation or above elbow | |
| | L6684 | | Y | Upper extremity addition, test socket, shoulder | |

| | Procedure | | T. | D 14 | Policy/ |
|------------|-----------|----|-----|---|--|
| Indicators | Code | PA | Lic | Description | Comments |
| | L6686 | | Y | disarticulation or interscapular thoracic Upper extremity addition, suction socket | |
| | L6687 | | Y | Upper extremity addition, frame type socket, below | |
| | | | | elbow or wrist disarticulation | |
| | L6688 | | Y | Upper extremity addition, frame type socket, above elbow or elbow disarticulation | |
| | L6689 | Y | Y | Upper extremity addition, frame type socket, shoulder disarticulation | |
| | L6690 | Y | Y | Upper extremity addition, frame type socket, interscapular-thoracic | |
| | L6691 | Y | Y | Upper extremity addition, removable insert, each | |
| | L6692 | Y | Y | Upper extremity addition, silicone gel insert or equal, each | |
| | L6693 | Y | Y | Upper extremity addition, external locking elbow, forearm counterbalance | |
| | L6694 | ¥ | Y | Addition to upper extremity prosthesis, below elbow/above elbow, custom fabricated from existing mold or prefabricated, socket insert, silicone gel, elastomeric or equal, for use with locking mechanism. | Limit to one per client per year without prior authorization. |
| | L6695 | | Y | Addition to upper extremity prosthesis, below elbow/above elbow, custom fabricated from existing mold or prefabricated, socket insert, silicone gel, elastomeric or equal, not for use with locking mechanism | |
| | L6696 | Y | Y | Addition to upper extremity prosthesis, below elbow/above elbow, custom fabricated socket insert for congenital or atypical traumatic amputee, silicone gel, elastomeric or equal, for use with or without locking mechanism, initial only (for other than initial, use code L6694 or L6695) | |
| | L6697 | | Y | Addition to upper extremity prosthesis, below elbow/above elbow, custom fabricated socket insert for other than congenital or atypical traumatic amputee, silicone gel, elastomeric or equal, for use with or without locking mechanism, initial only (for other than initial, use code L6694 or L6695) | |
| | L6698 | Y | Y | Addition to upper extremity prosthesis, below elbow/above elbow, lock mechanism, excludes socket insert | |
| | L6700 | | Y | Terminal device, hook, Dorrance or equal, model #3 | |
| | L6703 | Y | Y | Terminal device, passive hand/mitt, any material, any size | |
| | L6704 | Y | Y | Terminal device, sport/recreational/work attachment, any material, any size | |
| | L6706 | Y | Y | Terminal device, hook, mechanical, voluntary opening, any material, any size, lined or unlined | |

| | Procedure | | | | Policy/ |
|-------------------|-----------|----|-----|--|--------------------------------------|
| Indicators | | PA | Lic | Description | Comments |
| | L6707 | Y | Y | Terminal device, hook, mechanical, voluntary closing, any material, any size, lined or unlined | |
| | L6708 | Y | Y | Terminal device, hand, mechanical, voluntary opening, any material, any size | |
| | L6709 | Y | Y | Terminal device, hand, mechanical, voluntary closing, any material, any size | |
| | L6711 | Y | Y | Terminal device, hook, mechanical, voluntary opening, any material, any size, lined or unlined, pediatric. | Document Correction DC- 2009-1 |
| | L6712 | Y | Y | Teminal device, hook, mechanical, voluntary closing, any material, any size, lined or unlined, pediatric. | Document Correction DC- 2009-1 |
| | L6713 | Y | Y | Terminal device, hand, mechanical, voluntary opening, any material, any size, pediatric. | Document Correction DC- 2009-1 |
| | L6714 | Y | Y | Terminal device, hand, mechanical, voluntary closing, any material, any size, pediatric. | Document Correction DC- 2009-1 |
| | L6721 | Y | Y | Terminal device, hook or hand, heavy duty, mechanical, voluntary opening, any material, any size, lined or unlined. | Document Correction DC- 2009-1 |
| | L6722 | Y | Y | Terminal device, hook or hand, heavy duty, mechanical, voluntary closing, any material, any size, lined or unlined. | Document Correction DC- 2009-1 |
| | L6810 | Y | Y | Terminal device, pincher tool, Otto Bock or equal | |
| | L6881 | Y | Y | Automatic grasp feature, addition to upper limb prosthetic terminal device | |
| | L6882 | Y | Y | Microprocessor control feature, addition to upper limb prosthetic terminal device | |
| | L6883 | Y | Y | Replacement socket, below elbow/wrist disarticulation, molded to patient model, for use with or without external power | |
| | L6884 | Y | Y | Replacement socket, above elbow disarticulation, molded to patient model, for use with or without external power | |
| | L6885 | Y | Y | Replacement socket, shoulder disarticulation/interscapular thoracic, molded to patient model, for use with or without external power | |
| | L6890 | | Y | Terminal device, glove for above hands, production glove | |
| | L6895 | Y | Y | Terminal device, glove for above hands, custom glove | |
| | L6900 | Y | Y | Hand restoration (casts, shading and measurements included), partial hand, with glove, thumb or one finger remaining | |
| | L6905 | Y | Y | Hand restoration (casts, shading and measurements | |

| | Procedure | | | | Policy/ |
|------------|-----------|----|-----|--|----------|
| Indicators | Code | PA | Lic | Description | Comments |
| | | | | included), partial hand, with glove, multiple fingers remaining | |
| | L6910 | Y | Y | Hand restoration (casts, shading and measurements included), partial hand, with glove, no fingers remaining | |
| | L6915 | Y | Y | Hand restoration (shading and measurements included), replacement glove for above | |
| | L6920 | Y | Y | Wrist disarticulation, external power, self-suspended inner socket, removable forearm shell, Otto Bock or equal switch, cables, two batteries and one charger, switch control of terminal device | |
| | L6925 | Y | Y | Wrist disarticulation, external power, self-suspended inner socket, removable forearm shell, Otto Bock or equal electrodes, cables, two batteries and one charger, myoelectronic control of terminal device | |
| | L6930 | Y | Y | Below elbow, external power, self-suspended inner socket, removable forearm shell, Otto Bock or equal switch, cables, two batteries and one charger, switch control of terminal device | |
| | L6935 | Y | Y | Below elbow, external power, self-suspended inner socket, removable forearm shell, Otto Bock or equal electrodes, cables, two batteries and one charger, myoelectronic control of terminal device | |
| | L6940 | Y | Y | Elbow disarticulation, external power, molded inner socket, removable humeral shell, outside locking hinges, forearm, Otto Bock or equal switch, cables, two batteries and one charger, switch control of terminal device | |
| | L6945 | Y | Y | Elbow disarticulation, external power, molded inner socket, removable humeral shell, outside locking hinges, forearm, Otto Bock or equal electrodes, cables, two batteries and one charger, myoelectronic control of terminal device | |
| | L6950 | Y | Y | Above elbow, external power, molded inner socket, removable humeral shell, internal locking elbow, forearm, Otto Bock or equal switch, cables, two batteries and one charger, switch control of terminal device | |
| | L6955 | Y | Y | Above elbow, external power, molded inner socket, removable humeral shell, internal locking elbow, forearm, Otto Bock or equal, electrodes, cables, two batteries and one charger, myoelectronic control of terminal device | |
| | L6960 | Y | Y | Shoulder disarticulation, external power, molded inner | |

| Code Status | Procedure | | | | Daliay/ |
|-------------------|-----------|----|-----|---|---------------------|
| Indicators | Code | PA | Lic | Description | Policy/ Comments |
| | 0000 | | | socket, removable shoulder shell, shoulder bulkhead, | 00111110110 |
| | | | | humeral section, mechanical elbow, forearm, Otto | |
| | | | | Bock or equal switch, cables, two batteries and one | |
| | | | | charger, switch control of terminal device | |
| | L6965 | Y | Y | Shoulder disarticulation, external power, molded inner | |
| | | | | socket, removable shoulder shell, shoulder bulkhead, | |
| | | | | humeral section, mechanical elbow, forearm, Otto Bock or equal electrodes, cables, two batteries and one | |
| | | | | charger, myoelectronic control of terminal device | |
| | L6970 | Y | Y | Interscapular-thoracic, external power, molded inner | |
| | | | | socket, removable shoulder shell, shoulder bulkhead, | |
| | | | | humeral section, mechanical elbow, forearm, Otto | |
| | | | | Bock or equal switch, cables, two batteries and one | |
| | | | | charger, switch control of terminal device | |
| | L6975 | Y | Y | Interscapular-thoracic, external power, molded inner | |
| | | | | socket, removable shoulder shell, shoulder bulkhead, humeral section, mechanical elbow, forearm, Otto | |
| | | | | Bock or equal electrodes, cables, two batteries and one | |
| | | | | charger, myoelectronic control of terminal device | |
| | L7007 | Y | Y | Electric hand, switch or myoelectric controlled, adult | |
| | L7008 | Y | Y | Electric hand, switch or myoelectric, controlled, | |
| | | | | pediatric | |
| | L7009 | Y | Y | Electric hook, switch or myoelectric controlled, adult | |
| | L7040 | Y | Y | Prehensile actuator, Hosmer or equal, switch controlled | |
| | L7045 | Y | Y | Electronic hook, child, Michigan or equal, switch controlled | |
| | | | | | |
| | L7170 | Y | Y | Electronic elbow, Hosmer or equal, switch controlled | |
| | L7180 | Y | Y | Electronic elbow, Boston, Utah or equal, | |
| | | | | myoelectronically controlled | |
| | L7181 | Y | Y | Electronic elbow, microprocessor simultaneous | |
| | | | | control of elbow and terminal device | |
| | L7185 | Y | Y | Electronic elbow, adolescent, Variety Village or equal, | |
| | 17106 | 37 | Y | switch controlled | |
| | L7186 | Y | ĭ | Electronic elbow, child, Variety Village or equal, switch controlled | |
| | L7190 | Y | Y | Electronic elbow, adolescent, Variety Village or equal, | |
| | 2/1/0 | • | - | myoelectronically controlled | |
| | L7191 | Y | Y | Electronic elbow, child, Variety Village or equal, | |
| | | | | myoelectronically controlled | |
| | L7260 | Y | Y | Electronic wrist rotator, Otto Bock or equal | |
| | L7261 | Y | Y | Electronic wrist rotator, for Utah arm | |
| | L7266 | Y | Y | Servo control, Steeper or equal | |
| | L7272 | Y | Y | Analogue control, UNB or equal | |

| Code | | | | | |
|------------|-----------|----|-----|--|---------------------------------------|
| | Procedure | | | D | Policy/ |
| Indicators | | PA | Lic | Description | Comments |
| | L7274 | Y | Y | Proportional control, 6-12 volt, Liberty, Utah or equal | |
| | L7360 | Y | Y | Six volt battery, Otto Bock or equal, each | |
| | L7362 | Y | Y | Battery charger, six volt, each. | |
| | L7364 | Y | Y | Twelve volt battery, each | |
| | L7366 | Y | Y | Battery charger, twelve volt, each. | |
| | L7367 | Y | Y | Lithium ion battery, replacement | |
| | L7368 | Y | Y | Lithium ion battery charger | |
| | L7400 | Y | Y | Addition to upper extremity prosthesis, below elbow/wrist disarticulation, ultralight material (titanium, carbon fiber or equal) | |
| | L7401 | Y | Y | Addition to upper extremity prosthesis, above elbow disarticulation, ultralight material (titanium, carbon fiber or equal) | |
| | L7402 | Y | Y | Addition to upper extremity prosthesis, shoulder disarticulation/interscapular thoracic, ultralight material (titanium, carbon fiber or equal) | |
| | L7403 | Y | Y | Addition to upper extremity prosthesis, below elbow/wrist disarticulation, acrylic material | |
| | L7404 | Y | Y | Addition to upper extremity prosthesis, above elbow disarticulation, acrylic material | |
| | L7405 | Y | Y | Addition to upper extremity prosthesis, shoulder disarticulation/interscapular thoracic, acrylic material | |
| | L7499 | Y | Y | Upper extremity prosthesis, not otherwise specified | |
| # | L7500 | Y | Y | Repair of prosthetic device, hourly rate | |
| | L7510 | Y | Y | Repair prosthetic device, repair or replace minor parts | |
| | L7520 | Y | Y | Repair of prosthetic device, labor component, per 15 minutes | |
| | L7600 | Y | Y | Prosthetic donning sleeve, any material, each | |
| # | L7900 | | | Vacuum erection system | |
| | L8000 | | *** | Breast prosthesis, mastectomy bra | |
| | L8001 | | *** | Breast prosthesis, mastectomy bra, with integrated breast prosthesis form, unilateral | Not allowed with L8020 or L8030 |
| | L8002 | | *** | Breast prosthesis, mastectomy bra, with integrated breast prosthesis form, bilateral | Not allowed with L8020 or L8030 |
| | L8010 | | *** | Breast prosthesis, mastectomy sleeve | |
| | L8015 | | *** | External breast prosthesis garment, with mastectomy form, post mastectomy | |
| | L8020 | | *** | Breast prosthesis, mastectomy form | |
| | L8030 | | *** | Breast prosthesis, silicone or equal | |
| # | L8035 | | | Custom breast prosthesis, post mastectomy, molded to patient model | |
| | L8039 | Y | | Breast prosthesis, not otherwise specified | |

| Code | | | | | |
|----------------------|-------------------|----|-------|---|---------------------|
| Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
| # | L8040 | IA | Lic | Nasal prosthesis, provided by a non-physician | Comments |
| # | L8041 | | | Midfacial prosthesis, provided by a non-physician | |
| # | L8042 | | | Orbital prosthesis, provided by a non-physician | |
| # | L8042 | | | Upper facial prosthesis, provided by a non-physician | |
| # | L8043 | | | Hemi-facial prosthesis, provided by a non-physician | |
| # | L8044 | | | Auricular prosthesis, provided by a non-physician | |
| # | L8045 | | | Partial facial prosthesis, provided by a non-physician | |
| # | L8040 | | | | |
| # | L8047 | | | Nasal septal prosthesis, provided by a non-physician | |
| # | L8048 | | | Unspecified maxillofacial prosthesis, by report, | |
| | | | | provided by a non-physician | |
| # | L8049 | | | Repair or modification of maxillofacial prosthesis, | |
| | | | | labor component, 15 minute increments, provided by a | |
| | ¥ 0200 | | .111. | non-physician | |
| | L8300 | | *** | Truss, single with standard pad | |
| | L8310 | | *** | Truss, double with standard pads | |
| | L8320 | | *** | Truss, addition to standard pad, water pad | |
| | L8330 | | *** | Truss, addition to standard pad, scrotal pad | |
| | L8400 | | Y | Prosthetic sheath, below knee, each | |
| | L8410 | | Y | Prosthetic sheath, above knee, each | |
| | L8415 | | Y | Prosthetic sheath, upper limb, each | |
| | L8417 | | Y | Prosthetic sheath/sock, including a gel cushion layer, | |
| | | | | below knee or above knee, each | |
| | L8420 | | Y | Prosthetic sock, multiple ply, below knee, each | |
| | L8430 | | Y | Prosthetic sock, multiple ply, above knee, each | |
| | L8435 | | Y | Prosthetic sock, multiple ply, upper limb, each | |
| | L8440 | | Y | Prosthetic shrinker, below knee, each | |
| | L8460 | | Y | Prosthetic shrinker, above knee, each | |
| | L8465 | | Y | Prosthetic shrinker, upper limb, each | |
| | L8470 | | Y | Prosthetic sock, single ply, fitting, below knee, each | |
| | L8480 | | Y | Prosthetic sock, single ply, fitting, above knee, each | |
| | L8485 | | Y | Prosthetic sock, single ply, fitting, upper limb, each | |
| | L8499 | Y | Y | Unlisted procedure for miscellaneous prosthetic | |
| | | | | services | |
| # | L8500 | | | Artificial larynx, any type | |
| # | L8501 | | | Tracheostomy speaking valve | |
| # | L8505 | | | Artificial larynx replacement battery/accessory, any type | |
| # | L8507 | | | Tracheo-esophageal voice prosthesis, patient inserted, | |
| | | | | any type, each | |
| # | L8509 | | | Tracheao-esophogeal voice prosthesis, inserted by a | |
| | | | | licensed health care provider, any type | |
| # | L8510 | | | Voice amplifier | |
| # | L8511 | | | Insert for indwelling tracheoesophageal prosthesis, | |

| Code Status Indicators | Procedure Code | PA | Lic | Description | Policy/ Comments |
|------------------------------|-------------------|----|----------|---|----------------------|
| THOROUGH S | 0000 | | <u> </u> | with or without valve, replacement only, each | Comments |
| # | L8512 | | | Gelatin capsules or equivalent, for use with tracheoesophageal voice prosthesis, replacement only, per 10 | 1 unit = 10 capsules |
| # | L8513 | | | Cleaning device used with tracheoesophageal voice prosthesis, pipet, brush, or equal, replacement only, each | |
| # | L8514 | | | Tracheoesophageal puncture dilator, replacement only, each | |
| # | L8515 | | | Gelatin capsule, application device for use with tracheoesophageal voice prosthesis, each | |
| # | L8600 | | | Implantable breast prosthesis, silicone or equal | |
| # | L8603 | | | Injectable bulking agent, collagen implant, urinary tract, per 2.5 ml syringe, includes shipping and necessary supplies | 1 unit = 2.5 ml |
| # | L8606 | | | Injectable bulking agent, synthetic implant, urinary tract, 1 ml syringe, includes shipping and necessary supplies | 1 unit = 1 ml |
| # | L8609 | | | Artificial cornea | |
| # | L8610 | | | Ocular Implant | |
| # | L8612 | | | Aqueous shunt | |
| # | L8613 | | | Ossicular implant | |
| # | L8614 | | | Cochlear device/system | |
| # | L8615 | | | Headset/headpiece for use with cochlear implant device, replacement | |
| # | L8616 | | | Microphone for use with cochlear implant device, replacement | |
| # | L8617 | | | Transmitting coil for use with cochlear implant device, replacement | |
| # | L8618 | | | Transmitter cable for use with cochlear implant device, replacement | |
| # | L8619 | | | Cochlear implant external speech processor, replacement | |
| # | L8621 | | | Zinc air battery for use with cochlear implant device, replacement, each | |
| # | L8622 | | | Alkaline battery for use with cochlear implant device, any size, replacement, each | |
| # | L8623 | | | Lihium battery for use with cochlear implant device speech processor, other than ear level, replacement, each | |
| # | L8624 | | | Lihium battery for use with cochlear implant device speech processor, ear level replacement, each | |

| Code | | | | | |
|------------|-----------|----|------------|---|----------|
| | Procedure | | - . | 5 | Policy/ |
| Indicators | | PA | Lic | Description | Comments |
| # | L8630 | | | Metacarpophalangeal joint implant | |
| # | L8631 | | | Metacarpal phalangeal joint replacement, two or more | |
| | | | | pieces, metal(e.g., stainless steel or cobalt chrome), | |
| | | | | ceramic-like material (e.g., pyrocarbon), for surgical | |
| .,, | T 0 6 4 1 | | | implantation (all sizes, includes entire system) | |
| # | L8641 | | | Metatarsal joint implant | |
| # | L8642 | | | Hallux implant | |
| # | L8658 | | | Interphalangeal joint implant | |
| # | L8659 | | | Interphalangeal finger joint replacement, two or more | |
| | | | | pieces, metal (e.g., stainless steel or cobalt chrome), | |
| | | | | ceramic-like material (e.g., pyrocarbon) for surgical | |
| | | | | implantation, any size | |
| # | L8670 | | | Vascular graft material, synthetic, implant | |
| # | L8680 | | | Implantable neurostimulator electrode, each | |
| # | L8681 | | | Patient programmer (external) for use with | |
| | | | | implantable programmable neurostimulator pulse | |
| | | | | generator | |
| # | L8682 | | | Implantable neurostimulator radiofrequency receiver | |
| # | L8683 | | | Radiofrequency transmitter (external) for use with | |
| | | | | implantable neurostimulator radiofrequency receiver | |
| # | L8684 | | | Radiofrequency transmitter (external) for use with | |
| | | | | implantable sacral root neurostimulator receiver for | |
| .,, | T 0 60 5 | | | bowel and bladder management, replacement | |
| # | L8685 | | | Implantable neurostimulator pulse generator, single | |
| | T 0.00 | | | array, rechargeable, includes extension | |
| # | L8686 | | | Implantable neurostimulator pulse generator, single | |
| ш | 1.0607 | | | array, non-rechargeable, includes extension | |
| # | L8687 | | | Implantable neurostimulator pulse generator, dual | |
| # | 1 0/00 | | | array, rechargeable, includes extension | |
| # | L8688 | | | Implantable neurostimulator pulse generator, dual array, non-rechargeable, includes extension | |
| # | 1 0600 | | | | |
| # | L8689 | | | External recharging system for implanted neurostimulator, replacement only | |
| # | L8690 | | | Auditory osseointegrated device, includes all internal | |
| # | L8090 | | | | |
| # | L8691 | | | and external components | |
| # | L0091 | | | Auditory osseointegrated device, external sound processor, replacement | |
| # | L8695 | | | External recharging system for battery (external) for | |
| # | L0093 | | | use with implantable neurostimulator | |
| # | 1 0600 | | | - | |
| | L8699 | | | Prosthetic implant, not otherwise specified | |
| # | L9900 | | | Orthotic and prosthetic supply, accessory, and/or service component of another HCPCS L code | |
| | V2623 | | | Prosthetic eye, plastic, custom | |

Prosthetic and Orthotic Devices

| Code | | | | | |
|-------------------|-----------|----|-----|---|----------|
| | Procedure | | | | Policy/ |
| Indicators | Code | PA | Lic | Description | Comments |
| | V2624 | | | Polishing/resurfacing of ocular prosthesis | |
| | V2625 | | | Enlargement of ocular prosthesis | |
| | V2626 | | | Reduction of ocular prosthesis | |
| | V2627 | | | Scleral cover shell | |
| | V2628 | | | Fabrication and fitting of ocular conformer | |
| | V2629 | Y | | Prosthetic eye, other type | |

Provider Requirements

What is required from HRSA's P&O devices providers? (Refer to WAC 388-543-1200 [1])

HRSA requires a provider who supplies P&O devices and related supplies and services to an HRSA client to meet all of the following. The provider must:

- Have a proper business license;
- Have appropriately trained qualified staff;
- Be certified, licensed, and/or bonded, if required, to perform the services billed to HRSA. Out-of-state P&O providers must meet their state regulatory requirements; and
- Have an HRSA core provider agreement.

Who does HRSA reimburse for providing P&O devices and related supplies and services to HRSA clients?

(Refer to WAC 388-543-1200 [2])

HRSA may reimburse qualified providers for P&O devices, repairs, and related supplies and services on a fee-for-service (FFS) basis as follows:

- Licensed P&O providers who are licensed by the Washington State Department of Health (DOH) in P&O. This does not apply to medical equipment dealers and pharmacies that do not require licensure to provide selected P&O;
- All HCPCS codes with a "***" indicator in the licensure column may be provided by a supplier that has a DME or Pharmacy provider number as long as all other licensure requirements have been met.
- Physicians who provide medical equipment and supplies in the physician's office. HRSA
 may pay separately for medical supplies, subject to the provisions in HRSA's Physician'sRelated Services (RBRVS) fee schedule; and
- Out-of-state P&O providers who meet their state regulations.

Note: HRSA terminates from Medicaid participation any provider who violates program regulations and policies, as described in WAC 388-502-0030. (WAC 388-543-1200 [3])

What records must be kept? (Refer to WAC 388-502-0020)

Enrolled providers must:

- Keep legible, accurate, and complete charts and records to justify the services provided to each client, including, but not limited to:
 - ✓ Patient's name and date of birth;
 - ✓ Dates of service(s);
 - ✓ Name and title of person performing the service, if other than the billing practitioner;
 - ✓ Chief complaint or reason for each visit;
 - ✓ Pertinent medical history;
 - ✓ Pertinent findings on examination;
 - ✓ Medications, equipment, and/or supplies prescribed or provided;
 - ✓ Description of treatment (when applicable);
 - ✓ Recommendations for additional treatments, procedures, or consultations;
 - ✓ X-rays, tests, and results;
 - ✓ Plan of treatment and/or care, and outcome;
 - ✓ Specific claims and payments received for services; and
 - ✓ Any specifically required forms for the provision of P&O devices.
- Assure charts are authenticated by the person who gave the order, provided the care, or performed the observation, examination, assessment, treatment or other service to which the entry pertains.
- Make charts and records available to DSHS, its contractors, and the US Department of
 Health and Human Services, upon their request, <u>for at least six years from the date of
 service</u> or more if required by federal or state law or regulation.

Note: A provider may contact HRSA with questions regarding its programs. However, HRSA's response is based solely on the information provided to HRSA's representative at the time of inquiry, and in no way exempts a provider from following the laws and rules that govern HRSA's programs. (Refer to WAC 388-502-0020[2])

Authorization

What is prior authorization?

Prior authorization (PA) is HRSA's approval for certain medical services, equipment, or supplies, before the services are provided to clients, as a precondition for provider reimbursement. Expedited prior authorization (EPA) and limitation extensions (LE) are forms of prior authorization.

Is Prior Authorization required? [Refer to WAC 388-543-1600]

Yes! The Medical Assistance Administration (HRSA) requires prior authorization for certain purchases and repairs of medically necessary P&O devices and related supplies and services. Please refer to the PA column of the *Fee Schedule* (Section H) for items that require prior authorization.

HRSA bases its determination about which P&O devices and related supplies and services require PA or EPA on utilization criteria. HRSA considers all of the following when establishing utilization criteria:

- High cost;
- Potential for utilization abuse:
- Narrow therapeutic indication; and
- Safety.

How do I request prior authorization?

Providers must submit the request in writing to the Quality Utilization Section or call the authorization toll-free number at 1-800-292-8064. (See *Important Contacts* section.)

General Policies for Prior Authorization

[Refer to WAC 388-543-1800]

• For PA requests, HRSA requires the prescribing provider to furnish patient-specific justification for base equipment and each requested line item accessory or modification identified as a separate charge. HRSA does not accept general standards of care or industry standards for generalized equipment as justification.

- When HRSA receives an initial request for PA, the prescription(s) for those items or services cannot be older than three months from the date HRSA receives the request.
- HRSA requires certain information from providers to prior authorize the purchase of equipment. This information includes, but is not limited to, the following:
 - ✓ A detailed description of the item; and
 - ✓ Any modifications required, including the product or accessory number as shown in the manufacturer's catalog.
- HRSA prior authorizes By Report (BR) items that require PA and are listed in the *Fee Schedule* only if medical necessity is established and the provider furnishes all of the following information to HRSA:
 - ✓ A detailed description of the item or service to be provided;
 - ✓ The cost or charge for the item;
 - ✓ A copy of the manufacturer's invoice, price-list or catalog with the product description for the item being provided; and
 - ✓ A detailed explanation of how the requested item differs from an already existing code description.
- HRSA does not reimburse for purchase or repair of medical equipment that duplicates equipment the client already owns. If the provider makes such a request, HRSA requires the provider to submit a PA request and explain the following:
 - ✓ Why the existing equipment no longer meets the client's medical needs; or
 - ✓ Why the existing equipment could not be repaired or modified to meet those medical needs.
- A provider may resubmit a request for PA for an item or service that HRSA has denied. HRSA requires the provider to include new documentation that is relevant to the request.
- HRSA prior authorizes extensive maintenance that the manufacturer recommends be performed by an authorized dealer. HRSA requires the client to take responsibility for routine maintenance of a prosthetic or orthotic. If the client does not have the physical or mental ability to perform the task, HRSA requires the client's caregiver to be responsible. [WAC 388-543-2600 (4)]

Note: Written requests for prior authorization must be submitted to HRSA on a 1500 Claim Form with the date of service left blank and a copy of the prescription attached.

What is a limitation extension?

A limitation extension is when HRSA allows additional units of service for a client when the provider can verify that the additional units of service are medically necessary. Limitation extensions require prior authorization.

Note: Requests for limitation extensions must be appropriate to the client's eligibility and/or program limitations. Not all client eligibility groups cover all services.

How do I request a limitation extension?

In cases where the provider feels that additional services are medically necessary for the client, the provider must request pre-approval from HRSA in writing.

The request must state the following in writing:

- 1. The name and PIC number of the client;
- 2. The provider's name, provider number and fax number;
- 3. Additional service(s) requested;
- 4. Copy of last prescription and date dispensed;
- 5. The primary diagnosis code and HCPCS code or state assigned code; and
- 6. Client-specific clinical justification for additional services.

Send your request for a limitation extension to:

Division of Health Services Quality Support Quality Utilization Section Durable Medical Equipment/P&O Devices PO Box 45506 Olympia, WA 98504-5506 800.292.8064 360.586.5299 fax

What is expedited prior authorization?

The expedited prior authorization process (EPA) is designed to eliminate the need for written and telephonic requests for prior authorization for selected P&O device procedure codes. HRSA allows payment during a continuous 12-month period for this process.

To bill HRSA for P&O devices that meet the EPA criteria on the following pages, the provider must create a 9-digit EPA number. The first 6 digits of the EPA number must be **870000**. The last 3 digits must be the code number of the product and documented medical condition that meets the EPA criteria. Enter the EPA number on the 1500 Claim Form in the *Authorization Number* field or in the *Authorization* field when billing electronically.

Example: The 9-digit EPA number for purchase of a foot insert, removable, formed to patient foot for a client that meets one of the EPA criteria would be **870000780** (870000 = first 6 digits, 780 = product and documented medical condition).

Providers are reminded that EPA numbers are only for those products listed on the EPA Criteria Coding List. EPA numbers are not valid for:

- P&O devices requiring prior authorization through the P&O Devices program;
- Products for which the documented medical condition does not meet <u>all</u> of the specified EPA criteria; or
- Over-limitation requests.

The written or telephonic request process for PA must be used when a situation does not meet the criteria for EPA for a selected P&O device procedure code. Providers must submit the request in writing to the Quality Utilization Section or call the authorization toll-free number at 1-800-292-8064. (See *Important Contacts* section.) (**Refer to WAC 388-543-1900[3**])

Expedited Prior Authorization Guidelines:

- **A. Medical Justification (criteria)** All information must come from the client's prescribing physician or therapist, with an appropriately completed prescription. HRSA does not accept information obtained from the client or from someone on behalf of the client (e.g. family).
- **B. Documentation** The billing provider **must keep** documentation of the criteria in the client's file. Upon request, a provider must provide documentation to HRSA showing how the client's condition met the criteria for EPA. Keep documentation file for six (6) years. (**Refer to WAC 388-543-1900[4]**)

Note: HRSA may recoup any payment made to a provider under this section if the provider did not follow the expedited authorization process and criteria. Refer to WAC 388-502-0100. **(WAC 388-543-1900[5])**

EPA Criteria Coding Table

Prosthetics

| Procedure | EPA | | |
|----------------|------|---|--|
| Code | Code | Description | Criteria |
| L5683 L5681 | 787 | Addition to lower extremity, below knee/above knee, socket insert, suction suspension with or without locking mechanism | Initial purchase of one (1) L5683 and L5681 per initial, lower extremity prosthesis (one to wash, one to wear) allowed per 12-month period if any of the following criteria are met: 1) Short residual limb; 2) Diabetic; or 3) History of skin problems/open sores on stump |
| | | | Note: |
| | | | If the medical condition does not meet one of the above specified criteria, you must obtain prior authorization by submitting a request in writing to QUS (see Important Contacts) or by calling the authorization toll-free number at 1-800-292-8064. This EPA is allowed only one time per client, per 12-month period. It is the provider's responsibility to determine whether the EPA has been used for the client within 12 months prior to the provider's proposed date of service. EPA is for initial purchase only. It is not to be used for replacements of existing products. |

Orthotics

| Procedure | E+PA | | |
|----------------|------|--|---|
| Code | Code | Description | Criteria |
| L3030 | 780 | Foot insert, removable, formed to patient foot | One (1) pair allowed in a 12-month period if one of the following criteria is met: 1) Severe arthritis with pain; 2) Flat feet or pes planus with pain; 3) Valgus or varus deformity with pain; 4) Plantar fasciitis with pain; or 5) Pronation. |
| | | | Note: |
| | | | If the medical condition does not meet one of the above specified criteria, you must obtain prior authorization by submitting a request in writing to QUS (see Important Contacts) or by calling the authorization toll-free number at 1-800-292-8064. This EPA is allowed only one time per client, per 12-month period. It is the provider's responsibility to determine whether the EPA has been used for the client within 12 months prior to the provider's proposed date of service. |
| L3310 L3320 | 781 | Lift, elevation, heel & sole, per inch | For a client with a leg length discrepancy, allowed for as many inches as required (must be at least one inch), on one shoe per 12-month period. |

| Procedure | E+PA | | | | | | | |
|-----------|------|---------------------------------|--|--|--|--|--|--|
| Code | Code | Description | Criteria | | | | | |
| L3334 | 782 | Lift, elevation, heel, per inch | Allowed for as many inches as required (has to be at least one inch), for a client with a leg length discrepancy, on one shoe per 12-month period. Note: | | | | | |
| | | | - 1,000 | | | | | |
| | | | Lift is covered per inch, for no less than one (1) inch, for one shoe. For example: It is medically necessary for a client to have a two (2) inch lift for the left heel. Bill two units of L3334 using EPA # 870000782.* If the medical condition does not meet the above specified criteria, you must obtain | | | | | |
| | | | prior authorization by submitting a request in writing to QUS (see Important Contacts) or by calling the authorization toll-free number at 1-800-292-8064. 4) This EPA is allowed only one time per client, per 12-month period. It is the | | | | | |
| | | | provider's responsibility to determine whether the EPA has been used for the client within 12 months prior to the provider's proposed date of service. | | | | | |

*On-line clarification.

| Procedure | E+PA | | |
|-----------|------|---|---|
| Code | Code | Description | Criteria |
| L3000 | 784 | Foot insert, removable, molded to patient model, "UCB" type, Berkeley Shell, each | Purchase of one (1) pair per 12-month period for a client 16 years of age or younger allowed if any of the following criteria are met: 1) Required to prevent or correct pronation; 2) Required to promote proper foot alignment due to pronation; or 3) For ankle stability as required due to an existing medical condition such as hypotonia, Cerebral Palsy, etc. Note: 1) If the medical condition does not meet one of the above specified criteria, you must obtain prior authorization by submitting a request in writing to QUS (see <i>Important Contacts</i>) or by calling the authorization toll-free number at 800.292.8064. 2) This EPA is allowed only one time per client, per 12-month period. It is the provider's responsibility to determine whether the EPA has been used for the client within 12 months prior to the provider's proposed date of service. 3) If the client only medically requires one orthotic, right or left, prior authorization must be obtained. |

| Procedure | E+PA | | |
|----------------|------|--|--|
| Code | Code | Description | Criteria |
| L3215 L3219 | 785 | Orthopedic footwear, woman's or man's shoes, oxford. | Purchase of one (1) pair per 12-month period allowed if any of the following criteria are met: 1) When one or both shoes are attached to a brace; 2) When one or both shoes are required to accommodate a brace with the exception of L3030 foot inserts; 3) To accommodate a partial foot prosthesis; or 4) To accommodate club foot. |
| | | | Note: |
| | | | DSHS does not allow orthopedic footwear for the following reasons: a) To accommodate L3030 orthotics; b) Bunions; c) Hammer toes; d) Size difference (mismatched shoes); or e) Abnormal sized foot. DSHS only allows the following manufacturers of Orthopedic: a) Acor; b) Alden Shoe Company; c) Jerry Miller; d) Markell; e) P.W. Minor; f) Walkin-Comfort; and g) Hanger. If the medical condition does not meet one of the above specified criteria, you must obtain prior authorization by submitting a request in writing to QUS (see Important Contacts) or by calling the authorization toll-free number at 800.292.8064. EPA is allowed only one time per client, per 12-month period. It is the provider's responsibility to determine whether the client has already used all EPA in the period allowed under the EPA criteria. |

Prosthetic and Orthotic Devices

| Procedure | E+PA | | |
|-----------|------|---|--|
| Code | Code | Description | Criteria |
| L1945 | 786 | AFO, molded to patient model, plastic, rigid anterior tibial section (floor reaction) | Purchase of one per limb allowed per 12-month period if all of the following criteria are met: 1) Client is 16 years old or younger; and 2) Required due to a medical condition causing crouched gait. |
| | | | Note: |
| | | | If the medical condition does not meet one of the above specified criteria, you must obtain prior authorization by submitting a request in writing to QUS (see Important Contacts) or by calling the authorization toll-free number at 1-800-292-8064. EPA is allowed only one time per client, per 12-month period. It is the provider's responsibility to determine whether the client has already used all EPA in the period allowed under the EPA criteria. |

Reimbursement

General Reimbursement for P&O Devices and Related Supplies and Services (Refer to WAC 388-543-1400 and 388-543-2700)

- HRSA reimburses a qualified provider who serves a client who is not enrolled in a department-contracted managed care plan only when all of the following apply:
 - ✓ The provider meets all of the conditions in WAC 388-502-0100; and
 - ✓ HRSA does not include the item/service for which the provider is requesting reimbursement in other reimbursement rate methodologies. Other methodologies include, but are not limited to, the following:
 - ➤ Hospice providers' per diem reimbursement;
 - Hospital's diagnosis related group (DRG) reimbursement;
 - Managed care plans' capitation rate; and
 - Nursing facilities' per diem rate.
- A provider must not bill HRSA for the purchase of equipment supplied to the provider at no cost by suppliers/manufacturers.
- Reimbursement for P&O devices is limited to HCPCS/National Codes with the same level of coverage as Medicare.
- HRSA may adopt policies, procedure codes, and/or rates that are different than those set by Medicare, if HRSA determines that such actions are in the best interest of its clients.
- HRSA may pay for medical services rendered to a client only when HRSA is the payer of last resort.
- HRSA's maximum payment for medical equipment and supplies is the lesser of either of the following:
 - ✓ Provider's usual and customary charge; or
 - ✓ Established rates, unless the client is eligible for both Medicare and Medicaid (see *Billing* section).
- HRSA determines reimbursement for P&O devices according to a set fee schedule (see section H). HRSA considers Medicare's current fee schedule when determining maximum allowable fees. For By Report (BR) codes, HRSA reimburses 85% of the agreed upon fee.

- HRSA sets maximum allowable fees for P&O devices and related supplies and services using available published information, such as:
 - ✓ Commercial databases for price comparisons;
 - ✓ Manufacturers' catalogs;
 - ✓ Medicare fee schedules; and
 - ✓ Wholesale prices.
- HRSA evaluates and updates the maximum allowable fees for P&O devices at least once per year, independent of scheduled legislatively authorized vendor rate increases. Rates remain effective until the next rate change.

Specific Reimbursement for P&O Devices

(Refer to WAC 388-543-2700)

- HRSA's reimbursement for a P&O device includes the cost of any necessary molds.
- HRSA's hospital reimbursement rate includes any P&O devices required for surgery and/or placed during the hospital stay.
- Reimbursement for gender dysphoria surgery includes payment for all related prosthetics and supplies.

Purchased P&O Devices and Related Supplies

(Refer to WAC 388-543-1500)

- P&O devices and related supplies that HRSA purchases for a client are the client's property. HRSA reimbursement for covered P&O devices and related supplies includes all of the following:
 - Any adjustments or modifications to the equipment that are required within three months of the date of delivery. This does not apply to adjustments required because of changes in the client's medical condition;
 - ✓ Fitting and set-up; and
 - ✓ Instruction to the client or client's caregiver in the appropriate use of the equipment, device, and/or supplies.
- HRSA requires a provider to furnish to HRSA clients only new equipment that includes full manufacturer and dealer warranties.

- HRSA charges the dispensing provider for any costs it incurs to have another provider repair equipment if all of the following apply:
 - ✓ The dispensing provider is unwilling or unable to fulfill the warranty; and
 - ✓ The client still needs the equipment.
- HRSA rescinds purchase orders for the following reasons:
 - ✓ If the equipment was not delivered to the client before the client:
 - Dies:
 - Loses medical eligibility;
 - Becomes covered by a hospice agency; or
 - Becomes covered by an HRSA managed care plan.
 - A provider may incur extra costs for customized equipment that may not be easily resold. In these cases, for purchase orders rescinded per the stipulations listed above, HRSA may pay the provider an amount it considers appropriate to help defray these extra costs. HRSA requires the provider to submit justification sufficient to support such a claim.
 - ✓ A client may become a managed care plan client before HRSA completes the purchase of prescribed medical equipment. If this occurs:
 - HRSA rescinds the purchase order until the managed care primary care provider (PCP) evaluates the client; then
 - HRSA requires the PCP to write a new prescription if the PCP determines the equipment is still medically necessary (see *Definitions* section); then
 - The managed care plan's applicable reimbursement policies apply to the purchase or rental of the equipment.

Note: P&O devices placed during an inpatient hospital stay **are** included in the hospital reimbursement rate. HRSA does **not** reimburse separately under these circumstances.

Fee Schedule

You may view HRSA's Prosthetic and Orthotic Devices Fee Schedule on-line at

http://hrsa.dshs.wa.gov/RBRVS/Index.html

Understanding the fee schedule

• In the P.A. (Prior Authorization) column on the fee schedule:

Y means requires prior authorization; and

Y* means requires prior authorization only for clients 17 years of age and older.

• In the Licensure column on the fee schedule:

Y means licensure required; and

Y** means licensure required if prescribing treatment of scoliosis.

*** means the item can be provided by a DME or Pharmacy provider as long as other licensure requirements have been met.

- **HCPCS codes** that do not have a Medicaid Maximum Allowance established are listed as *By Report* (B.R.) or **Noncovered** (#).
- Please provide the following documentation for By Report procedures requiring prior approval:
 - (1) A detailed description of the item that will be provided.
 - (2) The procedure code that most closely describes the By Report item. If the item has been modified, note how that was done.
 - (3) If appropriate, the manufacturer's invoice, price list, a catalog with product description, and cost of itemized items.
- Modifiers:

RT = Right

LT = Left

RP = **Replacement**

Note: If dispensing new bilateral/single item(s), use modifiers RT, LT, as appropriate. If dispensing replacement for a previous prosthetic(s) or orthotic(s), use modifier RP.

Billing

What is the time limit for billing? [Refer to WAC 388-502-0150]

- HRSA requires providers to submit an initial claim, be assigned an internal control number (ICN), and adjust all claims in a timely manner. HRSA has two timeliness standards: 1) for initial claims; and 2) for resubmitted claims.
- The provider must submit claims as described in HRSA's billing instructions.
- HRSA requires providers to obtain an ICN for an **initial claim** within 365 days from any of the following:
 - ✓ The date the provider furnishes the service to the eligible client;
 - ✓ The date a final fair hearing decision is entered that impacts the particular claim;
 - ✓ The date a court orders HRSA to cover the services; or
 - ✓ The date DSHS certifies a client eligible under delayed¹ certification criteria.
- HRSA may grant exceptions to the 365 billing day time limit for **initial claims** when billing delays are caused by either of the following:
 - ✓ DSHS certification of a client for a retroactive² period; or
 - The provider proves to HRSA's satisfaction that there are other extenuating circumstances.

Delayed Certification - According to WAC 388-500-0005, delayed certification means department approval of a person's eligibility for a covered service made after the established application processing time limits. If, due to delayed certification, the client becomes eligible for a covered service that has already been provided, the provider must not bill, demand, collect, or accept payment from the client or anyone on the client's behalf for the service; and must promptly refund the total payment received from the client or anyone acting on the client's behalf and then bill HRSA for the service.

Eligibility Established After Date of Service but Within the Same Month - If the client becomes eligible for a covered service that has already been provided because the client applied to the department for medical services later in the same month the service was provided (and is made eligible from the first day of the month), the provider must not bill, demand, collect, or accept payment from the client or anyone acting on the client's behalf for the service; and must promptly refund the total payment received from the client or anyone acting on the client's behalf and then bill HRSA for the service.

Retroactive Certification - According to WAC 388-500-0005, retroactive period means the three calendar months before the month of application (month in which client applied). If, due to retroactive certification, the client becomes eligible for a covered service that has already been provided, the provider must not bill, demand, collect, or accept payment from the client or anyone acting on the client's behalf for any unpaid charges for the service; and may refund any payment already received from the client or anyone acting on the client's behalf, and after refunding the payment, the provider may bill HRSA for the service.

• Providers may **resubmit, modify, or adjust** any timely initial claim, <u>except prescription</u> drug claims, for a period of 36 months from the date of service. Prescription drug claims must be resubmitted, modified, or adjusted within 15 months from the date of service.

Note: HRSA does not accept any claim for resubmission, modification, or adjustment after the allotted time period listed above.

- The allotted time periods do not apply to overpayments that the provider must refund to DSHS. After the allotted time periods, a provider may not refund overpayments to HRSA by claim adjustment. The provider must refund overpayments to HRSA by a negotiable financial instrument such as a bank check.
- The provider, or any agent of the provider, must not bill a client or a client's estate when:
 - ✓ The provider fails to meet these listed requirements; and
 - ✓ HRSA does not pay the claim.

What fee should I bill HRSA for eligible clients?

Bill HRSA your usual and customary fee.

Exception: If billing Medicare Part B crossover claims, bill the amount submitted to Medicare.

How do I bill for services provided to Primary Care Case Management (PCCM) clients?

When billing for services provided to PCCM clients:

- Enter the referring physician or PCCM name in field 17 on the 1500 Claim Form; and
- Enter the seven-digit, HRSA-assigned identification number of the PCCM who referred the client for the service(s). If the client is enrolled with a PCCM and the PCCM referral number is not in field 17a when you bill HRSA, the claim will be denied.

How do I bill for clients who are eligible for Medicare and Medical Assistance?

If a client is eligible for both Medicare and Medical Assistance (otherwise known as "dualeligible"), **you must <u>first</u> submit a claim to Medicare and accept assignment within Medicare's time limitations**. HRSA may make an additional payment after Medicare reimburses you.

- If Medicare pays the claim, the provider must bill HRSA within six months of the date Medicare processes the claim.
- If Medicare denies payment of the claim, HRSA requires the provider to meet HRSA's initial 365-day requirement for initial claim (see page I.1).
- Codes billed to HRSA must match codes billed to Medicare when billed as a Medicare Part B crossover claim.

Medicare Part B

Benefits covered under Part B include: **Physician, outpatient hospital services, home health, durable medical equipment, and other medical services and supplies** not covered under Part A.

When the words "This information is being sent to either a private insurer or Medicaid fiscal agent," appear on your Medicare remittance notice, it means that your claim has been forwarded to HRSA or a private insurer for deductible and/or coinsurance processing.

If you have received a payment or denial from Medicare, but it does not appear on your HRSA Remittance Advice and Status Report (RA) within 45 days from Medicare's statement date, you should bill HRSA directly.

- If Medicare has made payment, and there is a balance due from HRSA, you must submit a 1500 Claim Form (with the "XO" indicator in field 19). Bill only those lines Medicare paid. Do not submit paid lines with denied lines. This could cause a delay in payment or a denial.
- If Medicare denies services, but HRSA covers them, you must bill on a 1500 Claim Form (without the "XO" indicator in field 19). Bill only those lines Medicare denied. Do not submit denied lines with paid lines. This could cause a delay in payment or a denial.
- If Medicare denies a service that requires prior authorization by HRSA, HRSA will waive the prior authorization requirement but will still require authorization. Authorization or denial of your request will be based upon medical necessity.

Note:

- ✓ Medicare/Medical Assistance billing claims must be received by HRSA within six (6) months of the Medicare EOMB paid date.
- ✓ A Medicare Remittance Notice or EOMB must be attached to each claim.

Payment Methodology – Part B

- MMIS compares HRSA's allowed amount to Medicare's allowed amount and selects the lesser of the two. (If there is no HRSA allowed amount, we use Medicare's allowed amount.)
- Medicare's payment is deducted from the amount selected above.
- If there is *no* balance due, the claim is denied because Medicare's payment exceeds HRSA's allowable.
- If there *is* a balance due, payment is made towards the deductible and/or coinsurance up to HRSA's maximum allowable.

HRSA cannot make direct payments to clients to cover the deductible and/or coinsurance amount of Part B Medicare. HRSA *can* pay these costs to the provider on behalf of the client when:

- 1) The provider accepts assignment; and
- 2) The total combined reimbursement to the provider from Medicare and Medicaid does not exceed Medicare or Medicaid's allowed amount, whichever is less.

Third-Party Liability

You must bill the insurance carrier(s) indicated on the client's Medical Identification card. An insurance carrier's time limit for claim submissions may be different from HRSA's. It is your responsibility to meet the insurance carrier's requirements relating to billing time limits, as well as HRSA's, prior to any payment by HRSA.

You must meet HRSA's 365-day billing time limit even if you haven't received notification of action from the insurance carrier. If your claim is denied due to any existing third-party liability, refer to the corresponding HRSA Remittance Advice and Status Report (RA) for insurance information appropriate for the date of service.

If you receive an insurance payment and the carrier pays you less than the maximum amount allowed by HRSA, or if you have reason to believe that HRSA may make an additional payment:

- Submit a completed claim form to HRSA;
- Attach the insurance carrier's statement or EOB;
- If rebilling, also attach a copy of the HRSA RA showing the previous denial; or
- If you are rebilling electronically, list the claim number (ICN) of the previous denial in the *Comments* field of the Electronic Media Claim (EMC).

Third-party carrier codes are available on HRSA's website at http://HRSA.dshs.wa.gov or by calling the Coordination of Benefits Section at 1-800-562-6136.

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| January 2007 | - G.6 - |

Completing the 1500 Claim Form

Attention! HRSA now accepts the new 1500 Claim Form.

- On November 1, 2006, HRSA began accepting the new 1500 Claim Form (version 08/05).
- **As of April 1, 2007**, HRSA will no longer accept the old HCFA-1500 Claim Form.

Note: HRSA encourages providers to make use of electronic billing options. For information about electronic billing, refer to the *Important Contacts* section.

Refer to HRSA's current *General Information Booklet* for instructions on completing the 1500 Claim Form. You may download this booklet from HRSA's website at: http://maa.dshs.wa.gov/download/Billing%20Instructions%20Web%20Pages/General%20Information.html or request a paper copy from the Department of Printing (see Important Contacts section).

The following 1500 Claim Form instructions relate to **Prosthetic and Orthotic Devices Billing Instructions**. Click the link above to view general 1500 Claim Form instructions.

For questions regarding claims information, call HRSA toll-free:

800.562.3022

1500 Claim Form Field Descriptions

| Field | Name | Field | Entry |
|-------|----------------|------------|--|
| No. | Name | Required | Entry |
| 21. | Diagnosis or | When | Enter the appropriate diagnosis code(s) in areas 1, 2, |
| | Nature of | applicable | 3, and 4. A valid ICD-9-CM code will be required. |
| | Illness or | | HRSA no longer allows the use of an unspecified/ |
| | Injury | | dummy diagnosis code such as V58.9. |
| | - - | | |

| 24B. | Place of Service | Yes | These are the only appropriate code(s) for this billing instruction: | | | |
|------|------------------|-----|--|--|--|--|
| | | | Code Number | To Be Used For | | |
| | | | 12 | Home | | |
| | | | 13 | Assisted Living Facility | | |
| | | | 31 | Nursing Facility | | |
| | | | 32 | Skilled Nursing Facility | | |
| | | | 99 | Other place of service | | |
| 24C. | Type of Service | No | | | | |
| 24E. | Diagnosis Code | Yes | Enter the ICD-9-CM diagnor procedure or service being in 24D). A diagnosis code is or line billed. Enter the code 9-CM. A valid ICD-9-CM colonger allows the use of an undiagnosis code such as V58. | billed (for each item listed s required for each service le exactly as shown in ICD-code is required. HRSA no unspecified/dummy 9. | | |
| 28. | Total Charge | Yes | Enter the sum of your chargor decimals in this field. HRSA does not accept "conclaim form must be totaled" | tinued" claim forms. Each | | |

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Common Questions Regarding Medicare Part B/ Medicaid Crossover Claims

Q: Why do I have to mark "XO" in black ink in box 19 on crossover claim?

A: The "XO" allows our mailroom staff to identify crossover claims easily, ensuring accurate processing for payment. Use black ink for the "XO" in box 19 on crossover claims.

Q: What fields do I use for 1500 Claim Form Medicare information?

| A: | In Field: | Please Enter: |
|-----------|-----------|---|
| | 19 | an "XO" |
| | 24K | Medicare's allowed charges |
| | 29 | Medicare's total deductible |
| | 30 | Medicare's total payment |
| | 32 | Medicare's EOMB process date, and the third-party |
| | | liability amount |

Q: When I bill Medicare denied lines to HRSA, why is the claim denied?

A: Your bill is not a crossover when Medicare denies your claim or if you are billing for Medicare-denied lines. The Medicare EOMB must be attached to the claim. Do not indicate "XO."

Q: How do my claims reach Medicaid after I've sent them to Medicare?

A: After Medicare has processed your claim, and if Medicare has allowed the services, in most cases Medicare will forward the claim to HRSA for any supplemental Medicaid payment. When the remarks code is, "MA07-The claim information has also been forwarded to Medicaid for review," it means that your claim has been forwarded to HRSA.

Q: What if my claim(s) does not appear on the RA?

A: If **Medicare has paid** and the Medicare crossover claim does not appear on the HRSA Remittance Advice and Status Report (RA) within 45 days of the Medicare statement date, you should bill HRSA the *paid lines* on the HCFA-1500 claim form **with** an "XO" in box 19.

If **Medicare denies** a service, bill HRSA the <u>denied lines</u>, using the HCFA-1500 claim form **without** an "XO" on the claim.

REMEMBER! Attach a copy of Medicare's EOMB.

REMEMBER! You must submit your claim to HRSA within six months of the Medicare statement date if Medicare has **paid** or 365 days from date of service if Medicare has **denied**.

Note: Claims billed to HRSA with payment by Medicare must be submitted with the same procedure code used to bill Medicare.

How to Complete the 1500 Claim Form for Medicare Part B/Medicaid **Crossovers**

The 1500 Claim Form, used for Medicare/Medicaid Benefits Coordination, cannot be billed electronically.

Attention! HRSA now accepts the new 1500 Claim Form (version 08/05). **As of April 1, 2007**, HRSA will no longer accept the old HCFA-1500 Claim Form.

Refer to HRSA's current General Information Booklet for instructions on completing the 1500 Claim Form. You may download this booklet from HRSA's website at: http://maa.dshs.wa.gov/download/Billing%20Instructions%20Web%20Pages/General%20Infor mation.html or request a paper copy from the Department of Printing (see Important Contacts section).

The following 1500 Claim Form instructions relate to **Prosthetic and Orthotic Devices Billing Instructions**. Click the link above to view general 1500 Claim Form instructions.

> For questions regarding claims information, call HRSA toll-free: 800.562.3022

1500 Claim Form Field Descriptions

| Field No. | Name | Field Required | Entry |
|--------------|--------------|-------------------|--|
| 19. | Reserved For | Yes | When Medicare allows services, enter XO to indicate |
| | Local Use | | this is a crossover claim. |
| 22. | Medicaid | When | If this billing is being resubmitted more than six (6) |
| | Resubmission | Applicable | months from Medicare's paid date, enter the |
| | | | Internal Control Number (ICN) that verifies that |
| | | | your claim was originally submitted within the time |
| | | | limit. [The ICN number is the claim number listed |
| | | | on the Remittance Advice and Status Report (RA).] |
| | | | Also enter the three-digit denial Explanation of |
| | | | Benefits (EOB). |

| Field No. | Name | Field Required | Entry | | | | | | |
|--------------|----------------------|-------------------|---|-------------------------------------|--|--|--|--|--|
| 24B. | Place of Service | Yes | These are the only appro | priate code(s) for this | | | | | |
| | | | billing instruction: | | | | | | |
| | | | Code Number | To Be Used For | | | | | |
| | | | 12 | Home | | | | | |
| | | | 13 | Assisted Living Facility | | | | | |
| | | | 31 | Nursing Facility | | | | | |
| | | | 32 | Skilled Nursing Facility | | | | | |
| | | | 99 | Other place of service | | | | | |
| 24C. | Type of Service | No | | | | | | | |
| 24K. | Reserved for | Yes | Use this field to show Me | dicare allowed charges. | | | | | |
| | Local Use | | Enter the Medicare allow | ed charge on each detail | | | | | |
| | | | line of the claim (see sam | ple). | | | | | |
| 27. | Accept Assignment | Yes | Check yes. | | | | | | |
| 29. | Amount Paid | Yes | Enter the Medicare Dedu | Medicare Deductible here. Enter the | | | | | |
| | | | amount as shown on Medicare's Remittance Notice | | | | | | |
| | | | and Explanation of Medi | care Benefits (EOMB). If | | | | | |
| | | | you have more than six (| 6) detail lines to submit, | | | | | |
| | | | please use multiple 1500 | Claim Forms (see field 24) | | | | | |
| | | | and calculate the deducti | ble based on the lines on | | | | | |
| | | | each form. Do not includ | e coinsurance here. | | | | | |
| 30. | Balance Due | Yes | Enter the Medicare Total | Payment. Enter the | | | | | |
| | | | amount as shown on Med | licare's Remittance Notice | | | | | |
| | | | or Explanation of Medica | are Benefits (EOMB). If | | | | | |
| | | | you have more than six (| 6) detail lines to submit, | | | | | |
| | | | please use multiple HCFA | A claim forms (see field 24) | | | | | |
| | | | and calculate the Medica | re payment based on the | | | | | |
| | | | lines on each form. Do no | ot include coinsurance here. | | | | | |
| 32. | Name and | Yes | Enter Medicare Statemen | nt Date and any Third- | | | | | |
| | Address of | | Party Liability Dollar An | nount (e.g., auto, employee- | | | | | |
| | Facility Where | | sponsored, supplemental | insurance) here, if any. If | | | | | |
| | Services Are | | there is insurance payme | nt on the claim, you must | | | | | |
| | Rendered | | also attach the insurance | Explanation of Benefits | | | | | |
| | | | (EOB). Do not include co | insurance here. | | | | | |

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| 2. PATIENTS | NAME (L | ast Nar | me, First | Name, | Middle | nitial) | | 3. P | ATIENT 2 0 | SBIRT | DATE | si | X | 4. INSURED'S | NAME (| Last Na | me, Firs | t Name | , Middle | Initial) | |
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| 5. PATIENTS 100 AN | | | | | | | | | | | ONSHIP: | | | 7. INSURED'S | ADORE | SS (No. | , Street | | | | |
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| 12. PATIENT | S OR AU | | | | | | | ING A SI | | | | normation | necessary | 13. INSURED | | | | | | TURE I author ysician or supp | |
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