



UnitedHealthcare® West Benefit Interpretation Policy

# Durable Medical Equipment (DME), Prosthetics, Corrective Appliances/Orthotics (Non-Foot Orthotics) and Medical Supplies Grid

Policy Number: BIP051.Z Effective Date: May 1, 2023

Instructions for Use

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#### **Related Benefit Interpretation Policy**

<u>Durable Medical Equipment (DME), Prosthetics, Corrective Appliances/</u>
 Orthotics (Non Foot Orthotics) and Medical Supplies

### **Medical Supplies Grid**

Note: This policy is based, in part, upon Medicare DME MAC and/or Medicare criteria

It	em	Coverage	Comments
Abdominal binder	Surgical	Medical Supply*	Only when used as a dressing/holder; also see <u>Dressings, Surgical</u> .
	Non-surgical	Corrective Appliance/ Orthotic	<ul> <li>Covered when all of the following criteria are met:</li> <li>Serves a medical purpose and it is only associated with treating an illness, injury or malformed body member;</li> <li>Provides support and counter force (a force in a defined direction of a magnitude at least as great as a rigid or semi-rigid support) on the limb or body part that is being used to brace;</li> <li>Not used to supply compression therapy (e.g., to reduce size, volume, or swelling of a body member or to help circulation)</li> <li>Not used for convenience or appearance;</li> <li>Not used for cosmetic purposes.</li> </ul>
Aero Chamber (spa	cer)	Not Covered	Not Covered as DME; may be available as a pharmacy benefit

ltem	Coverage	Comments
Air Conditioner/Air Cleaner/Purifier/ Electrostatic Machines or other Environmental Equipment	Not Covered	EOC Exclusion. Environmental control, not primarily medical in nature
Air-fluidized Bed (Bead), e.g., Clinitron	DME	Home use of an air-fluidized bed is recommended when all of the following criteria are met:  The member has a stage 3 (full thickness tissue loss) or stage 4 (deep tissue destruction) pressure sore  The member is bedridden or chair bound as a result of severely limited mobility  The member would require institutionalization in the absence of an air-fluidized bed  The air-fluidized bed is ordered in writing by the member's attending physician based upon a comprehensive assessment and evaluation of the member after completion of a course of conservative treatment designed to optimize conditions that promote wound healing  The conservative treatment course must have been at least one month in duration without progression toward wound healing. The month of conservative treatment may include some period in an institution as long as there is documentation available to verify that the necessary conservative treatment has been rendered  Conservative treatment must include:  Frequent repositioning of the member with particular attention to relief of pressure over bony prominences (usually every 2 hours)  Use of a specialized support surface (Group 2) designed to reduce pressure and shear forces on healing ulcers and to prevent new ulcer formation  Necessary treatment to resolve any wound infection  Optimization of nutrition status to promote wound healing  Debridement by any means (including wet to dry dressings, which does not require an occlusive covering) to remove devitalized tissue from the wound bed  Maintenance of a clean, moist bed of granulation tissue with appropriate moist dressings protected by an occlusive covering, while the wound heals  A trained adult caregiver is available to assist the member with activities of daily living, fluid balance, dry skin care, repositioning, recognition and management of altered status, dietary needs, prescribed treatments, and management and support of the air-fluidized bed system and its problems, such as leakage  A physician directs the home treatment regimen and re-evalu
		The member requires treatment with wet soaks or moist wound dressings that are not protected with an impervious covering, such as plastic wrap or other occlusive material

Item	Coverage	Comments
		<ul> <li>The caregiver is unwilling or unable to provide the type of care required by the member on an air-fluidized bed</li> <li>Structural support is inadequate to support the weight of the air-fluidized bed system, which generally weighs 1,600 pounds or more</li> <li>Electrical system is insufficient for the anticipated increase in energy consumption</li> </ul>
		Coverage criteria apply; refer to the NCD for Air-Fluidized Bed (280.8). (Accessed March 1, 2022)
Air Splint	Medical Supply*	Clear plastic splints inflated by air used temporarily on fractured, broken, crushed or burned limbs
Alternating Pressure Pads, Gel Flotation Devices, Lamb's Wool Pads/Sheep Skins (Group 1 pressure reducing support surfaces)	DME	<ul> <li>Covered if the member meets:</li> <li>Criterion 1, or</li> <li>Criteria 2 or 3; and</li> <li>At least one of criteria 4-7.</li> <li>1. Criteria</li> <li>2. Completely immobile - i.e., member cannot make changes in body position without assistance</li> <li>3. Limited mobility - i.e., member cannot independently make changes in body position significant enough to alleviate pressure</li> <li>4. Any stage pressure ulcer on the trunk or pelvis</li> <li>5. Impaired nutritional status</li> <li>6. Fecal or urinary incontinence</li> <li>7. Altered sensory perception</li> <li>8. Compromised circulatory status</li> </ul>
Alternating Pressure Pads, Low Air Loss or Powered Flotation without Low Air Loss (Group 2 pressure reducing support surfaces)	DME	<ul> <li>Mattresses (Pressure Reducing) are recommended if the member meets the following:</li> <li>Criterion a and b and c, or</li> <li>Criterion d, or</li> <li>Criterion e and f.</li> </ul> Criteria 1. Multiple stage II pressure ulcers (see Appendix I for details) located on the trunk or pelvis 2. Member has been on a comprehensive ulcer treatment program for at least the past month, which has included the use of an appropriate Group 1 support surface. The comprehensive treatment should include the following: <ul> <li>a. Education of the member and caregiver on the prevention and/or management of pressure ulcers</li> <li>b. Regular assessment by a nurse, physician, or other licensed healthcare practitioner (usually at least weekly for a member with a stage III or IV ulcer)</li> </ul>

ltem	Coverage	Comments
		<ul> <li>c. Appropriate turning and positioning</li> <li>d. Appropriate wound care (for a stage II, III, or IV ulcer)</li> <li>e. Appropriate management of moisture/incontinence</li> <li>f. Nutritional assessment and intervention consistent with the overall plan of care</li> <li>3. The ulcers have worsened or remained the same over the past month.</li> <li>4. Large or multiple stage III or IV pressure ulcer(s) on the trunk or pelvis</li> <li>5. Recent myocutaneous flap or skin graft for a pressure ulcer on the trunk or pelvis (surgery within the past 60 days)</li> <li>6. The member has been on a Group 2 or 3 support surface immediately prior to a recent discharge from a hospital or nursing facility (discharge within the past 30 days)</li> <li>Note: If the member is using a Pressure Reducing Mattress, there should be a care plan established by the physician or home care nurse, which includes the elements listed above. The support surface provided for the member should be one in which the member does not "bottom out." Bottoming out is the finding that an outstretched hand can readily palpate the bony prominence (coccyx or lateral trochanter) when it is placed palm up beneath the undersurface of the mattress or overlay and in an area under the bony prominence. The bottoming out criterion should be tested with the member in the supine position with their head flat, in the supine position with their head slightly elevated (no more than 30 degrees), and in the side lying position.</li> <li>Note: When a Pressure Reducing Mattress is provided following a myocutaneous flap or skin graft, recommendation generally is limited to 60 days from the date of surgery.</li> </ul>
		Continued Use  Continued use of a Pressure Reducing Mattress is recommended until the ulcer is healed or, if healing does not continue, there is documentation in the medical record to show the following:  Other aspects of the care plan are being modified to promote healing, or  The use of the Pressure Reducing Mattress is medically necessary for wound management
Ambulatory Boot	Corrective Appliance/ Orthotic	Also known as surgical boot
Ambulatory Cardiac Event Monitoring (example: Holter Monitor, Event Monit Patch-Type Monitor, Zio Patch)	Medical or, Supply*	Refer to the Medical Management Guideline (MMG) titled Cardiac Event Monitoring.
Ankle-Foot Non-ambulatory Orthosis (AFO)/ Static or dynamic positioning ankle foot orthoses (AFO)	- Orthotic	Static or dynamic positioning ankle-foot orthoses (AFO) is covered when criteria are met. Refer to the DME MAC <u>LCD for Ankle-Foot/Knee-Ankle-Foot Orthoses (L33686)</u> . (Accessed March 1, 2022)

Item		Coverage	Comments
			<ol> <li>Covered if either all of criteria 1-4 or criterion 5 is met:</li> <li>Plantar flexion contracture of the ankle with a dorsiflexion on passive range of motion testing of at least 10 degrees</li> <li>Reasonable expectation of the ability to correct the contracture</li> <li>Contracture is interfering or expected to interfere significantly with the member's functional abilities</li> <li>Used as a component of a therapy program which includes active stretching of the involved muscles and/or tendons.</li> <li>Member has plantar fasciitis</li> </ol>
	mbulatory Prop Splint	Not Medically Necessary	A foot drop splint/recumbent positioning device and replacement interface will be denied as not medically necessary in a member with foot drop who is nonambulatory because there are other more appropriate treatment modalities.
Or Kr Fo (K Ar (e. wa pn	atory nkle-Foot rthosis (AFO) nee-Ankle- not Orthosis AFO)/ mbulatory ng., cam neumatic olint	Corrective Appliance/ Orthotic	Ankle-foot orthoses (AFO) and knee-ankle-foot orthoses (KAFO) are covered when criteria are met.  Refer to the DME MAC LCD for Ankle-Foot/Knee-Ankle-Foot Orthoses (L33686).  (Accessed March 2, 2022)  Ankle-foot orthoses (AFO) are covered for ambulatory members with weakness or deformity of the foot and ankle, which require stabilization for medical reasons, and have the potential to benefit functionally.  Knee-ankle-foot orthoses (KAFO) are covered for ambulatory member's for whom an ankle-foot orthosis is covered and for whom additional knee stability is required.  AFO, KAFO, and braces for ankle, foot, and knee used solely for athletic sports are not covered.  AFOs and KAFOs that are molded-to-member model are covered for ambulatory member's when the basic coverage criteria listed above are met and one of the following criteria are met:  The member could not be fit with a prefabricated AFO, or  The condition necessitating the orthosis is expected to be permanent or of long standing duration (more than 6 months), or  There is a need to control the knee, ankle or foot in more than 1 plane, or  There is a documented neurological, circulatory, or orthopedic status that requires custom fabricating over a model to prevent tissue injury, or  The member has a healing fracture which lacks normal anatomical integrity or anthropometric proportions.
Apnea Monitor (Infant or Ch	nild)	DME	There must be documentation of sleep apnea by a sleep study and a history of apnea events. Rental only. Not covered for adults.
Artificial Eye		Prosthetic	Covered for member with absence or shrinkage of an eye due to birth defect, trauma or surgical removal. Coverage includes polishing and resurfacing. Orbital implants are reimbursed as surgical implants.

ŀ	tem	Coverage	Comments
Artificial Larynx or	ficial Larynx or Electronic Speech Aid	Prosthetic	Coverage for member post laryngectomy or permanently inoperative larynx condition; disposable aid Not Covered.
			There are two types of speech aids. One operates by placing a vibrating head against the throat; the other amplifies sound waves through a tube which is inserted into the user's mouth. A member who has had radical neck surgery and/or extensive radiation to the anterior part of the neck would generally be able to use only the "oral tube" model or one of the more sensitive and more expensive "throat contact" devices.
Artificial Limbs – Lower Limb	Standard	Prosthetic	Covered when medical criteria are met; Refer to the InterQual® Client Defined CP: Durable Medical Equipment, Prosthetic Devices, Wigs, Specialized, Microprocessor or Myoelectric Upper Limbs (Custom) – UHG Click <a href="here">here</a> to view the InterQual® criteria.
	Bionic		Note: Members may have coverage for this item under the prosthetic benefit in some plans. Refer to the
	C-leg (microprocessor- controlled knee- shin system)		member's EOC/SOB or contact the Customer Service Department to determine coverage eligibility.
	Myoelectric		
Artificial Limbs – Upper Limb	Standard	Prosthetic	<ul> <li>A determination of the medical necessity for the prosthesis is based on the member's potential functional abilities. Potential function ability is based on the reasonable expectations of the prosthetist and treating physician, considering factors including, but not limited to, the following:</li> <li>The member's past history (including prior prosthetic use if applicable); and</li> <li>The member's current condition including the status of the residual limb and the nature of the other medical problems; and</li> <li>The member's desire to use a prosthesis.</li> </ul>
			Body Powered Prostheses – Upper Limb
			Upper limb functional body-powered prostheses are powered and controlled by gross body movements, a harness, and cable system. The following are basic requirements necessary for a member to be a candidate for a body-powered prosthesis:  Sufficient residual limb length Sufficient musculature Sufficient range of motion
			A member must possess at least one more of the following gross body movements to be able to control a body-powered prosthesis:  Glenohumeral flexion Scapular abduction or adduction Chest expansion

It	em	Coverage	Comments
			Shoulder depression and elevation
	Myoelectric		<b>Note</b> : Members may have coverage for this item under the prosthetic benefit in some plans. Refer to the member's EOC/SOB or contact the Customer Service Department to determine coverage eligibility
Back Brace			See <u>Spinal Orthosis</u> .
Back Support (post	cure chair)	Not Covered	Not primarily medical in nature
Bath Accessories	Bath Tub Lifts and Seats	Not Covered	Not primarily medical in nature
	Transfer Bench	Not Covered	Not primarily medical in nature
Beds and	Hospital, fixed	DME	Member must meet one or more of the following criteria.
Accessories	height		<ul> <li>Requires positioning of the body in ways not feasible with an ordinary bed. Elevation of the head/upper body less than 30 degrees does not usually require the use of a hospital bed</li> <li>Require positioning of the body in ways not feasible with an ordinary bed, for alleviation of pain</li> <li>Require the head of the bed to be elevated more than 30 degrees most of the time due to congestive heart failure, chronic pulmonary disease or problems with aspiration (pillows or wedges should be considered first)</li> <li>Require traction equipment that can only be attached to a hospital bed</li> </ul>
			NCD for Hospital Beds (280.7) (Accessed April 6, 2022)
	Hospital, variable height	DME	Member must meet one of the criteria for the fixed height bed (as listed above) and must require a bed height different than a fixed height bed in order to permit transfer to a chair, wheelchair or standing position;
			and
			<ul> <li>Variable height feature of a hospital bed is covered for one of the following conditions:</li> <li>Severe arthritis and other injuries to lower extremities; e.g., fractured hip. The condition requires the variable height feature to assist the member to ambulate by enabling the member to place his or her feet on the floor while sitting on the edge of the bed;</li> <li>Severe cardiac conditions. For those cardiac members who are able to leave bed, but who must avoid the strain of "jumping" up or down;</li> <li>Spinal cord injuries, including quadriplegic and paraplegic member's multiple limb amputee and stroke members. For those members who are able to transfer from bed to a wheelchair, with or without help; or</li> <li>Other severely debilitating diseases and conditions, if the variable height feature is required to assist the member to ambulate.</li> </ul>
	Hospital, semi- electric	DME	Member must meet one of the criteria for the fixed height bed (as listed above) and must require frequent or immediate changes in body position.

Ite	em	Coverage	Comments
	Hospital, total electric	Not Covered	The electric height adjustment feature is a convenient item therefore does not meet the definition of DME.
	Hospital, heavy duty extra wide	DME	Recommended for member's meeting criteria for a fixed height hospital bed and the member's weight is more than 350 pounds, but does not exceed 600 pounds.
	Hospital, extra heavy duty	DME	Recommended for member's meeting criteria for a fixed height hospital bed and the member's weight exceeds 600 pounds.
	Lounge (power or manual)	Not Covered	Not primarily medical in nature
	Mattress	DME	Only when part of a hospital bed
	Oscillating	Not Covered	Institutional equipment; inappropriate for home use. Does not meet the definition of DME.
	Over Bed Tables	Not Covered	Not primarily medical in nature
	Pressure Reducing	DME	<ul> <li>Mattresses (Pressure Reducing) are recommended if the member meets the following:</li> <li>Criteria 1 and 2 and 3, or</li> <li>Criterion 4, or</li> <li>Criteria 5 and 6.</li> </ul>
			<ol> <li>Multiple stage II pressure ulcers (see Appendix I for details) located on the trunk or pelvis</li> <li>Member has been on a comprehensive ulcer treatment program for at least the past month, which has included the use of an appropriate Group 1 support surface. The comprehensive treatment should include the following:         <ol> <li>Education of the member and caregiver on the prevention and/or management of pressure ulcers</li> <li>Regular assessment by a nurse, physician, or other licensed healthcare practitioner (usually at least weekly for a member with a stage III or IV ulcer)</li> <li>Appropriate turning and positioning</li> <li>Appropriate wound care (for a stage II, III, or IV ulcer)</li> <li>Appropriate management of moisture/incontinence</li> <li>Nutritional assessment and intervention consistent with the overall plan of care</li> </ol> </li> <li>The ulcers have worsened or remained the same over the past month.</li> <li>Large or multiple stage III or IV pressure ulcer(s) on the trunk or pelvis</li> <li>Recent myocutaneous flap or skin graft for a pressure ulcer on the trunk or pelvis (surgery within the past 60 days)</li> <li>The member has been on a Group 2 or 3 support surface immediately prior to a recent discharge from a hospital or nursing facility (discharge within the past 30 days)</li> </ol>

Iten	n	Coverage	Comments
			Note: If the member is using a Pressure Reducing Mattress, there should be a care plan established by the physician or home care nurse, which includes the elements listed above. The support surface provided for the member should be one in which the member does not "bottom out." Bottoming out is the finding that an outstretched hand can readily palpate the bony prominence (coccyx or lateral trochanter) when it is placed palm up beneath the undersurface of the mattress or overlay and in an area under the bony prominence. The bottoming out criterion should be tested with the member in the supine position with their head flat, in the supine position with their head slightly elevated (no more than 30 degrees), and in the side lying position.  Note: When a Pressure Reducing Mattress is provided following a myocutaneous flap or skin graft, recommendation generally is limited to 60 days from the date of surgery.  Continued Use  Continued Use  Continued use of a Pressure Reducing Mattress IS recommended until the ulcer is healed or, if healing does not continue, there is documentation in the medical record to show the following:  Other aspects of the care plan are being modified to promote healing, or
			The use of the Pressure Reducing Mattress is medically necessary for wound management.
	Side rails	DME	Only if part of hospital bed and member's condition requires bed side rails
Bed Baths (home type	e)	Not Covered	Not primarily medical in nature
Bed Board		Not Covered	Not primarily medical in nature
Bed Lifter (bed elevate	or)	Not Covered	Not primarily medical in nature
Bed Cradle		DME	Covered when necessary to prevent contact with the bed coverings.
Bed Pan (autoclavable	e, hospital type)	DME	If member is bed bound
Bed Specs (prism glas	sses)	Not Covered	Not primarily medical in nature
Bed Wetting Alarms		Not Covered	Not primarily medical in nature
Bilevel Positive Airway	y Pressure (BiPAP)	DME	Covered as DME.
Bili-lights/Bili-blankets	s (phototherapy)	DME	Covered when medically necessary.
Blood Glucose Analyz Colorimeter	zer-reflectance	Not Covered	Unsuitable for home use. Does not meet the definition of DME.
Blood Pressure Monitors Sphygmomanometer	· ·	DME	Only for members on home dialysis; fully and semi-automatic (member activated) portable monitors are not covered.
Bone Stimulator also I Osteogenic Stimulator Ultrasonic)		DME	Criteria apply; refer to the MMG titled <u>Electrical and Ultrasound Bone Growth Stimulators</u> .

	Item	Coverage	Comments
Braces		Corrective Appliance/ Orthotic	Excludes orthodontic braces; also see <u>AFO/KAFO</u> or <u>Knee Orthosis</u> or <u>Spinal Orthosis</u> (body jacket) or <u>Back Brace</u> .
Braille Teaching 1	Гехt	Not Covered	Educational, not primarily medical in nature
Bras (post-surger	y)	Prosthetic	Two covered initially, with replacements thereafter due to normal wear and tear; coverage includes custom fittings.
Breast Prosthesis	(external)	Prosthetic	Covered for members who have had a mastectomy or lumpectomy. Refer to the Benefit Interpretation Policy (BIP) titled Post Mastectomy Surgery.  Initial prosthesis is covered for the useful lifetime of the prosthesis, with replacements thereafter due to normal wear and tear. Replacement of the same type is covered at any time when it's lost or irreparably damaged.  The useful lifetime expectancy for silicone breast prostheses is 2 years. The useful lifetime expectancy for nipple prosthesis is 3 months. For fabric, foam, or fiber filled breast prostheses, the useful lifetime expectancy is 6 months. Replacement sooner than the useful lifetime because of ordinary wear and tear
D 16 11 0		DNAF	will be denied as noncovered.
Counseling	ipport, Supplies and	DME	Refer to the MMG titled <u>Preventive Care Services</u> .
Cam Walkers (als Boot)	o known as Walking		See Ankle Foot Orthosis (AFO)/Knee Ankle Foot Orthosis (KAFO)
Canes	Quad or straight	DME	Covered when patient meets the Mobility Assistive Equipment clinical criteria.  Refer to the  NCD for Durable Medical Equipment Reference List (280.1)  NCD for Mobility Assistive Equipment (280.3)  DME MAC LCD for Canes and Crutches (L33733)  (Accessed March 2, 2022)
	White	Not Covered	Not primarily medical in nature
Car Seats		Not Covered	Not primarily medical in nature
Casts (e.g., plaste	er, fiberglass)	Medical Supply*	Used to reduce fractures or dislocations.
Catheters and Supplies	Closed Drainage Bags	Prosthetic (Covered as MPE)	Only for members with nonfunctioning bladder or permanent incontinence as medically required

Item	Coverage	Comments
External Urinary Collection Devices (e.g., male external catheters and female pouches/meatal cups)	Prosthetic (Covered as MPE)	Only for members with nonfunctioning bladder or permanent incontinence when used as an alternative to an indwelling catheter. Male external catheters are limited to no more than 35 per month and female external urinary collection devices are limited to no more than one meatal cup per week or one pouch per day. Requests for a greater quantity must be documented by a participating physician as medically necessary.
Foley/ Indwelling	Prosthetic (Covered as MPE)	Only for members with nonfunctioning bladder or permanent incontinence as medically required.  Limited to no more than one catheter per month for routine catheter maintenance. Requests for a greater quantity must be documented by a participating physician as medically necessary.
Intermittent Urinary Catheters	Prosthetic (Covered as MPE)	Intermittent catheterization is covered when basic coverage criteria are met and the member or caregiver can perform the procedure.  When clean, non-sterile catheterization technique is used, replacement of intermittent catheters on a weekly basis is covered unless there is documentation of the medical necessity for more frequent replacement. Non-sterile lubricating gel is covered for up to 8 units of service (8 oz.) per month. An individual packet of lubricant is not covered.  Intermittent catheterization using sterile technique is covered when the member requires catheterization and the member meets one of the following criteria (1-5):  1. The member resides in a nursing facility 2. The member is immunosuppressed (e.g., on a regimen of immunosuppressive drugs post-transplant) 3. The member has radiologically documented vesico-ureteral reflux while on a program of intermittent catheterization 4. The member is a spinal-cord injured female with neurogenic bladder who is pregnant (for duration of pregnancy only) 5. The member has had distinct, recurrent urinary tract infections, while on a program of clean intermittent catheterization, twice within the 12-month prior to the initiation of sterile intermittent catheterization  Note: The medical necessity for use of sterile intermittent catheterization for reasons other than the criteria (1-5) listed above may be presented for individual consideration.  For each episode of covered sterile catheterization, one catheter and an individual packet of lubricant or an intermittent catheter kit are covered.
Leg Drainage Bags	Prosthetic (Covered as MPE)	Only for members with nonfunctioning bladder or permanent incontinence who are ambulatory or are chair or wheelchair bound.

ltem		Coverage	Comments
Cervical Collar	Semi-rigid	Corrective Appliance/ Orthotic	Provides increased cervical support over foam.
	Soft	Corrective Appliance/ Orthotic	Minimal cervical support for sprains/strains
	Rigid	Corrective Appliance/ Orthotic	Covered post-surgery.
Cervical Pillow		Not Covered	Not primarily medical in nature
Cervical Thoracic L Orthoses (CTLSO)	umbar Sacral		See <u>Spinal Orthosis</u> ; also see <u>Scoliosis Orthosis</u> .
Chair (adjustable)		DME	Only for members on home dialysis; refer to the BIP titled <u>Dialysis Services</u> .
Chemical Test Strip	os		Refer to the BIP titled <u>Diabetic Management</u> , <u>Services and Supplies</u> .
Clavicle Support/S	Clavicle Support/Splint		Used to keep the clavicle in position following acute injury or post-operative care.
Cleft Palate Prosthe	esis	Prosthetic	Only for cleft lip and palate deformities as a result of congenital malformation
Cochlear Implant (E of Device)	Cochlear Implant (External Component of Device)		Considered as high-end prosthetic device.  Criteria apply; refer to the BIP titled <u>Hearing Services</u> .  Also refer to the MMG titled <u>Cochlear Implants</u> .
Water circulating	<ul> <li>Cold Therapy</li> <li>Cold Packs /Cool Jackets</li> <li>Water circulating cold pad with pump (e.g., Polar Units)</li> </ul>		Not medically necessary. Alternative therapy available with the same outcomes.
Collagen Implant		Prosthetic	Covered as prosthetic.
Colostomy Bag		Prosthetic	See Ostomy Supplies item.
Commode (without wheels only)	Bedside	DME	Covered when member is physically incapable of utilizing regular toilet facilities. This would occur when (1) member is confined to a single room, or (2) member is confined to one level of the home environment and there is not toilet on that level, or (3) member is confined to the home and there are no toilet facilities in the home.
	Chair Foot Rest	Not Covered	
	Elevated Seat (raised toilet seat)	Not Covered	

Item	Coverage	Comments
Communication Devices (e.g., computers, personal digital assistants, speech generating devices)  Except artificial larynxes	Not Covered	EOC Exclusion
Contact Lens, Hydrophilic Soft (external)		Covered under the medical benefit. Coverage criteria apply. Refer to the BIP titled <u>Vision Care and Services</u> .
Continuous Passive Motion (CPM)	DME	Covered for member's who have received a total knee replacement. To qualify for coverage, use of the device must commence within 2 days following surgery. In addition, coverage is limited to that portion of the 3-week period following surgery during which the device is used in the member's home. There is insufficient evidence to justify coverage of these devices for longer periods of time or for other applications.
Continuous Positive Airway Pressure (CPAP)	DME	Covered as DME.  Ventilators must <b>not</b> be billed using codes for CPAP (E0601) or bi-level PAP (E0470, E0471, E0472).  Using the CPAP or bi-level PAP HCPCS codes to bill a ventilator is incorrect coding, even if the ventilator is only being used in CPAP or bi-level mode. Ventilators, such as the Trilogy Device, should not be used for obstructive sleep apnea or in place of a Respiratory Assist Device.  Refer to Correct Coding and Coverage of Ventilators.
Corset	Corrective Appliance/ Orthotic	Covered as Corrective Appliance/ Orthotic.
Cough Assist Device	DME	<ul> <li>Mechanical in-exsufflation devices are covered for member's who meet both of the following criteria:</li> <li>They have a neuromuscular disease, and</li> <li>This condition is causing a significant impairment of chest wall and/or diaphragmatic movement, such that it results in an inability to clear retained secretions.</li> </ul>
Cranial Band / Helmet(Cranial Orthosis)	Corrective Appliance/ Orthotic	Refer to the MMG titled <u>Plagiocephaly and Craniosynostosis Treatment</u>
Crutches, Crutch Tips and Handles	DME	Covered when patient meets the Mobility Assistive Equipment clinical criteria.  Refer to the:  NCD for Mobility Assistive Equipment (MAE) (280.3)  DME MAC LCD for Canes and Crutches (L33733)  Also refer to the Coverage Summary titled Mobility Assistive Equipment (MAE).  Note: Crutch substitute (HCPCS code E0118) is not covered. There is insufficient published clinical literature demonstrating safety and effectiveness in the Medicare population to establish the medical necessity for this device. Refer to the:

ltem		Coverage	Comments
			CGS News & Publication-E0118 – Crutch Substitute
			Noridian Article E0118 - Crutch Substitute
Dehumidifier		Not Covered	Environmental control, not primarily medical in nature
Diabetic Supplies (lancets, injection ai	0.0		Refer to the BIP titled <u>Diabetic Management</u> , <u>Services and Supplies</u> .
Dental Splint			See <u>Splints</u> .
Dialysis Home Kit, F	Peritoneal	DME	Only for members on home dialysis
Diapers		Not Covered	Hygienic supplies, non-reusable
Disposable Sheets		Not Covered	Hygienic item; non-reusable disposable supplies
Dressings/ Bandages	Non-surgical Dressings/ Bandages (e.g., Ace bandages)	Medical Supply*	Only when provided in the physician's office, otherwise considered over the counter
	Surgical Dressings	Medical Supply DME Prosthetic	Surgical dressings are limited to primary dressings (therapeutic or protective coverings applied directly to a wound) or secondary dressings (dressings that serve a therapeutic or protective function and are needed to secure a primary dressing, e.g., tape, roll gauze, transparent film) that are medically necessary for the treatment of a wound caused by, or treated by, a surgical procedure or wound debridement.  Surgical dressings may be covered as:  Medical supply when provided the physician's office.  DME when ordered by the treating physician or other healthcare professional for the member's home use in conjunction with a durable medical equipment (e.g., infusion pumps).  Prosthetic when ordered by the treating physician or other healthcare professional for the member's home use as dressing for surgical wound or for wound debridement or in conjunction with a prosthetic device (e.g., tracheostomy).
Easy Stand/Tilt Sta	nd	Not Covered	Not primarily medical in nature
Egg Crate		Not Covered	See Alternating Pads.
Elbow Orthosis		Corrective Appliance	Used for compression of tissue or to limit motion. Custom molded covered only when member cannot be fitted with a prefabricated elbow support.
Electrical Stimulation	H-wave Stimulation Device	Not Medically Necessary	Insufficient clinical evidence supporting effectiveness.
Devices (for chronic pain)	Interferential Device	Not Medically Necessary	Refer to the MMG titled <u>Electrical Stimulation for the Treatment of Pain and Muscle Rehabilitation</u> .
	Tens Unit	DME	Transcutaneous electrical nerve stimulator (TENS) are covered when coverage criteria are met. Refer to the NCD for Transcutaneous Electrical Nerve Stimulator (TENS) for Acute Post-operative Pain (10.2).

It	em	Coverage	Comments
			Local Coverage Determinations (LCDs)/Local Coverage Articles (LCAs) exist and compliance with these policies is required where applicable. Refer to the DME MAC <u>LCD for Transcutaneous Electrical Nerve Stimulators (TENS) (L33802)</u> .
			For coverage of supplies necessary for TENS; refer to the <u>NCD for Supplies Used in the Delivery of Transcutaneous Electrical Nerve Stimulation (TENS) and Neuromuscular Electrical Stimulation (NMES) (160.13)</u> .
Electrical Stimulation (Neuromuscular, N		DME	Coverage criteria apply; refer to the MMG titled Electrical Stimulation for the <u>Treatment of Pain and Muscle Rehabilitation</u> .
Electrical Stimulation Electromagnetic The Healing or Diathern (e.g., Diapulse)	erapy for Wound	Not Covered	Refer to MMG Electrical Stimulation and Electromagnetic Therapy for Wounds.
Electronic Speech	Aids		See Artificial Larynx.
Electric Tumor Trea	atment Field Therapy		Refer to the MMG titled Electric Tumor Treatment Field Therapy.
Elevators		Not Covered	Not primarily medical in nature
Emesis Basin		Not Covered	Not primarily medical in nature
Enuresis Training It	em (penile clamp)	Prosthetic	For members with urinary incontinence
Esophageal Dilator		Not Covered	Physician instrument, not appropriate for home use
Exercise Equipmentypes of bicycles)	t (e.g., barbells, all	Not Covered	Not primarily medical in nature
Face Masks	Oxygen	DME	Covered if member is on oxygen.
	Surgical	Not Covered	Non-reusable disposable items
Facial Prosthesis		Prosthetic	Facial prostheses are covered when there is a loss or absence of facial tissue due to disease, trauma, surgery, or a congenital defect.
Fluidic Breathing A	Fluidic Breathing Assister		See IPPB Machines.
Flutter Device/ Oscillatory Positive Expiratory Pressure Devices		DME	Covered as DME.
Foot Cradle	Foot Cradle		See Bed Cradle.
Foot Orthotic		Corrective Appliance/ Orthotic	For diabetics only when criteria are met; Refer to the BIP titled Shoes and Foot Orthotics.

ltem	Coverage	Comments
Formula (enteral feedings)		Coverage criteria apply. Refer to the BIP titled <i>Enteral and Oral Nutritional Therapy</i> for <u>OK Members</u> , <u>OR Members</u> , <u>TX Members</u> , and <u>WA Members</u> .  Also refer to the BIP titled <u>Home Health Care</u> .  Also see <u>Pumps</u> .
Gait Belt	Not Covered	Does not meet the definition of DME.
Gait Trainers		A gait trainer (or sometimes referred to as a rollator) is a term used to describe certain devices that are used to support a patient during ambulation. Gait trainers are billed using one of the codes for walkers. If a gait trainer has a feature described by one of the walker attachment codes (E0154 - E0157) that code may be separately billed. Other unique features of gait trainers are not separately payable and may not be billed with code E1399. If a supplier chooses to bill separately for a feature of a gait trainer that is not described by a specific HCPCS code, then code A9900 must be used.  Refer to the DME MAC LCD for Walkers (L33791). (Accessed April 11, 2022)
Grab Bars (for bath and toilet)	Not Covered	Not primarily medical in nature
Gradient Pressure Stockings (e.g., Jobst stockings)		See Stockings.
Hearing Aid		Refer to the BIP titled <u>Hearing Services</u> . Also refer to the MMG titled <u>Hearing Aids and Devices Including Wearable</u> , <u>Bone-Anchored and Semi-Implantable</u> .
Heat Lamp	Not Covered	Not primarily medical in nature
Heater (portable)	Not Covered	Not primarily medical in nature
Heating and Cooling Plants	Not Covered	Not primarily medical in nature
Helmets (Safety Equipment)	Not Covered	Not primarily medical in nature
Heparin/ saline flushes	DME	Covered if member meets the homebound status and heparin flush is necessary to maintain patency of the line.
High Frequency Chest Wall Compression Devices (e.g., ThAIRapy® vest)	DME	Refer to the MMG titled Airway Clearance Devices.
Home Prothrombin Time International Normalized Ratio (INR) Monitoring)/ Coagulation Monitor	DME	Refer to the NCD for Home Prothrombin Time/International Normalized Ratio (PT/INR) Monitoring for Anticoagulation Management (190.11) for more detailed benefit information. (This NCD is distinct from, and makes no changes to the clinical laboratory NCD for Prothrombin Time (PT) (190.17))  Also refer to the Medicare Benefit Policy Manual, Chapter 15, §60.1 – Incident To Physician's Professional Services.

Ite	em	Coverage	Comments
			<ul> <li>Notes:</li> <li>Test materials continue to include 4 tests. Frequency of reporting requirements shall remain the same.</li> <li>Home INR monitoring is not covered for members with porcine valves unless covered by local Medicare contractors.</li> <li>Refer to the Medicare Claims Processing Manual, Chapter 32, Section 60 – Coverage and Billing for Home Prothrombin Time (PT/INR) Monitoring for Home Anticoagulation Management</li> </ul>
Humidifier	For use with C-PAP or BiPAP (heated or non-heated)	DME	Covered as DME.
	For use with Respiratory Assist Devices		For coverage criteria for Respiratory Assist Devices, see Respiratory Assist Devices.
	For use with Oxygen System	DME	Covered if criteria for oxygen are met.
	Room or Central Heating System Types	Not Covered	Environmental control equipment; not medical in nature
Hydraulic Lifts			See <u>Lifts</u> .
Hypothermic Blanke	ets	Not Covered	Not primarily medical in nature
Immobilizer (extrem	ity)		See Knee Orthosis.
Incontinence Contro (mechanical and hy		Prosthetic	For members with permanent anatomic and neurologic dysfunction of the bladder
Incontinence Pads		Not Covered	Non-reusable disposable items
Infusion Pump		DME	See <u>Pumps</u> .
Inhalation Machine			See Nebulizers, or Humidifiers, or IPPB Machines.
Injectors, Jet pressu	Injectors, Jet pressure powered injectors		Refer to the BIP titled <u>Diabetic Management</u> , <u>Services and Supplies</u> .
Insulin pump, including insulin and necessary supplies		DME	Criteria apply; refer to the BIP titled <u>Diabetic Management</u> , <u>Services and Supplies</u> ,  Also refer to the MMG titled <u>Continuous Glucose Monitoring and Insulin Delivery for Managing Diabetes</u> .
IV Pole (Intravenous	IV Pole (Intravenous)		Covered if medically necessary.
Intermittent Positive (IPPB) Machines	Pressure Breathing	DME	When breathing is severely impaired (includes fluidic breathing assisters)
Jacuzzi		Not Covered	Not primarily medical in nature

	Item	Coverage	Comments
Knee Orthosis (e.g., knee immobilizer, range of motion knee orthosis, rigid ace design knee orthosis)		Corrective Appliance/ Orthotic	Custom molded covered when member cannot be fitted with prefabricated immobilizer.
Lamb's Wool Pads	S/Sheep Skins		See Alternating Pressure Pads.
Leotard (pressure	garment)	Not Covered	Not primarily medical in nature, therefore, does not meet definition of DME.
Lifts	Bathtub or Toilet	Not Covered	Not primarily medical in nature
	Hydraulic (Hoyer)	DME	Covered if the member's condition is such that periodic movement is necessary to effect improvement or to arrest or retard deterioration in the member's condition.
	Motorized (electric), Ceiling Modified	Not Covered	
	Seat Lift Mechanism	DME	<ul> <li>Covered when criteria are met in the NCD.</li> <li>Notes:</li> <li>Coverage is limited to the seat lift mechanism and installation of the mechanism only. Other related items and services such as costs for the chair or chair upholstery are not covered.</li> <li>Lift mechanism which operates by spring release with a sudden, catapult-like motion and jolts the patient from a seated to a standing position is not covered.</li> <li>Refer to the NCD for Seat Lift (280.4)</li> </ul>
	For Wheelchairs/ Scooters/ POVs	Not Covered	Not primarily medical in nature
	Trunk/Vehicle Modification	Not Covered	Not primarily medical in nature
Light Therapy Box Box)	(Therapeutic Light	Not Covered	Not primarily medical in nature; therefore does not meet the definition of DME.  For the treatment of psoriasis, see <u>Ultraviolet Cabinet</u> .
Lumbar-sacral (LSO)		Corrective Appliance/ Orthotic	See <u>Spinal Orthosis</u> ; also see <u>Scoliosis Orthosis</u> .
Lymphedema Pum nonsegmental)	Lymphedema Pumps (segmental and nonsegmental)		Coverage criteria apply; refer to the MMG titled <u>Pneumatic Compression Devices</u> .  Note: Complex Decongestive Physiotherapy/CDP is considered a medical treatment rather than part of rehabilitation/therapy, therefore, CDP is not subject to rehabilitation/therapy co-payment nor benefit maximum.

	Item	Coverage	Comments
Lymphedema Sleeve/ Compression Garments/Bandages (Wrap)		Prosthetic DME	Covered if Medically Necessary.  Note: Complex Decongestive Physiotherapy/CDP is considered a medical treatment rather than part of rehabilitation/therapy, therefore, CDP is not subject to rehabilitation/therapy copayment nor benefit maximum.  Reference  Women's Health and Cancer Rights Act of 1998
Mandibular Device	e (for sleep apnea)	DME	Covered when medical criteria are met.  Refer to the MMG titled Obstructive and Central Sleep Apnea Treatment.
Massage Devices		Not Covered	Not primarily medical in nature
Mattress			See <u>Beds</u> .
Mobile Stander		Not Covered	See standing frames
	Small Volume, electric	DME	Covered for medications approved for delivery by a nebulizer, including nebulized medications for asthma.  Also may be covered when it is medically necessary to administer appropriate inhalation medications for the management of COPD, cystic fibrosis, HIV, pneumocystosis, complications of organ transplants or thick or tenacious pulmonary secretions.
	Large Volume, Non-Disposable	DME	When medically necessary to deliver humidity to a member with thick, tenacious secretions, who has cystic fibrosis, bronchiectasis, a tracheostomy, or a tracheobronchial stent; Not Covered when used predominantly to provide room humidification.  Also covered when medically necessary to deliver humidity to a member with thick, tenacious secretions, who has cystic fibrosis, bronchiectasis, a tracheostomy, or a tracheobronchial stent. Not covered when used predominantly to provide room humidification.
	Large Volume, Disposable	Not Covered	Acceptable alternative available; Not primarily medical in nature; Disposable items are not considered DME by definition.
	Ultrasonic	Not Covered	Offers no proven clinical advantage over a standard nebulizer.
	Portable (AC/DC)	DME	Only one nebulizer is allowed for in home use when Medically Necessary. (Stationary/Portable). Nebulizers are not allowed for out of home use as it does not meet definition of DME.
	Medication		Covered through the member's supplemental pharmacy benefit when listed in the formulary.
Negative Pressure	Negative Pressure Wound Therapy Pump		See <u>Vacuum Assisted Closure Device</u> .
Neuromuscular El (NMES)	ectrical Stimulator		See <u>Electrical Stimulation Devices</u> .

	ltem	Coverage	Comments
Noncontact Non T Therapy	hermal Wound		Refer to the MMG titled <u>Noncontact Warming Therapy</u> , <u>Ultrasound Therapy and Fluorescence Imaging for Wounds</u> .
Nutritional	Enteral		Refer to the BIPs titled Enteral and Oral Nutritional Therapy for OK Members, OR Members, TX
Therapy	Parenteral	DME	Members, and WA Members, and Parenteral Nutrition Therapy.  Also refer to BIP Home Health Care.  Also see Pumps.
Obturator, palatal		Prosthetic	Only for surgically acquired deformity or trauma; Used to replace or fill in a missing palate or portion of the palate; Includes the denture when the denture or a portion of denture is an integral part (built-in) of the obturator.
Orthopedic Shoes			See <u>Shoes</u> .
Ostomy Supplies		Prosthetic	Supplies include but are not limited to irrigation/flushing equipment and other supplies directly related to care of the member's ostomy.
Oxygen equipment and	Stationary	DME	Covered when medical criteria are met; Refer to the InterQual® CP: Durable Medical Equipment, Home Oxygen Therapy. Click

It	em	Coverage	Comments
Paraffin Bath Unit	Portable	DME	Covered when the member has undergone a successful trial period of paraffin therapy ordered by a physician and the member's condition is expected to be relieved by a long term use of this modality.
	Standard	Not Covered	Not appropriate for home use
Parallel Bars		Not Covered	Support exercise equipment; Primarily for institutional use
Peak Expiratory Flo	w Meter, hand-held	DME	For the self-monitoring of member's with pure asthma when used as part of a comprehensive asthma management program
Percussor (Non- Vest type)	Electric or pneumatic, home model	DME	Covered for mobilizing respiratory tract secretions in member's with chronic obstructive lung disease, chronic bronchitis or emphysema, when member or operator of powered percussor has received appropriate training by a physician or therapist, and no one competent to administer manual therapy is available.
			For ThAIRapy® Vest System, Refer to the MMG titled Airway Clearance Devices.
	Intrapulmonary Percussive Ventilator (IPV)	Not Covered	No data to support the effectiveness of the device in the home setting; Inappropriate for home use; Therefore does not meet the definition of DME.
Personal or Comfor	t Items	Not Covered	Not primarily medical in nature
Pessary	Pessary		For prolapse of the uterus or nonsurgical treatment of rectocele and cystocele
Pleurx bottles and t	ubing	DME	Covered as DME.
Pneumatic Compre	ssion Devices	DME	Criteria apply; refer to the MMG titled Pneumatic Compression Devices.
Pneumatic Splints			See Ankle Foot Orthosis (AFO)/Knee-Ankle-Foot Orthosis (KAFO).
Porcine (Pig) Skin [	Dressings	Medical Supply*	When used as an airtight (occlusive) dressing for burns, donor sites, bedsores (decubiti), and ulcers/wounds
Postural Drainage E	Boards	DME	For members with chronic pulmonary condition
Power Operated Ve (POV)/Scooters	Power Operated Vehicles (POV)/Scooters		See Wheelchairs.
Power traction equipment/devices (e.g., VAX-D®, DRX9000, SpineMED™, Spina System™, Lordex® Decompression Unit, DRS System™)			See <u>Traction</u> .
Pulse Oximeter		DME	A pulse oximeter is covered in the home to monitor oxygen saturation when medically necessary for the safe management of a medical diagnosis that would otherwise require treatment at a higher level of care. Coverage requires that an individual be available in the home with the requisite training to interpret and apply pulse oximetry data. Medical conditions that may warrant a home pulse oximeter include, but are not limited to:

Item		Coverage	Comments
			<ul> <li>Infants with severe chronic lung disease, such as bronchopulmonary dysplasia</li> <li>Premature infants being actively monitored for apnea</li> <li>Members who require mechanical home ventilation</li> <li>Members being weaned off of a home ventilator or oxygen therapy</li> <li>Members with a tracheostomy who require tracheal suctioning to manage pulmonary secretions</li> <li>Members with a severe cardiopulmonary diagnosis that requires immediate adjustment of oxygen flow rates</li> <li>Members with a progressive neuromuscular condition that impairs the muscles of respiration (e.g., ALS, Muscular Dystrophy, Myasthenia Gravis)</li> <li>The use of a home pulse oximeter to monitor oxygen saturations in COPD or Asthma is not considered medically necessary in the absence of special circumstances such as those outlined above.</li> </ul>
Pumps, including medications and necessary supplies	Enteral		Refer to the BIP titled <i>Enteral and Oral Nutritional Therapy</i> for <u>OK Members</u> , <u>OR Members</u> , <u>TX Members</u> , and <u>WA Members</u> .  Pumps, tubing and supplies necessary to deliver the enteral formula is covered.  Also refer to the BIP titled <u>Home Health Care</u> .
	Infusion	DME	Covered when medical criteria are met; Refer to the InterQual® Medicare: Durable Medical Equipment, External Infusion Pumps. Click <a href="here">here</a> to view the InterQual® criteria.  External infusion pumps for vancomycin (J3370) are Not Covered.  Note: There is insufficient evidence to support the necessity of using an external infusion pump, instead of a disposable elastomeric pump or the gravity drip method, to administer vancomycin in a safe and appropriate manner.  Implantable infusion pumps for infusion of heparin in the treatment of recurrent thromboembolic disease are not covered.
	Insulin, external	DME	Refer to the BIP titled Diabetic Management, Services and Supplies.
	Insulin, implantable	Not Covered	
	Pain Control	DME	Covered when medical criteria are met; Refer to the InterQual® Medicare: Durable Medical Equipment, External Infusion Pumps. Click <a href="here">here</a> to view the InterQual® criteria.
	Parenteral	DME	Refer to the BIP titled Parenteral Nutrition Therapy.
	For Erectile Dysfunction		See <u>Vacuum Pump</u> .
Punctal Plug	Punctal Plug		For treatment of dry eyes
Purewick Urine Collection System			Refer to the MMG titled Omnibus Codes.

ltem		Coverage	Comments
Ramp (wheelchair)	Ramp (wheelchair)		Not primarily medical in nature
Recliner (chair)	Recliner (chair)		Member must be on home dialysis.
Respiratory Assist [	Respiratory Assist Devices		Coverage criteria apply; see Humidifier. Also refer to the MMG titled Obstructive and Central Sleep <u>Apnea Treatment</u> .
Rib Belt		Corrective Appliance/ Orthotic	<ul> <li>Covered when all of the following criteria are met:</li> <li>Serves a medical purpose and it is only associated with treating an illness, injury or malformed body member</li> <li>Provides support and counter force (a force in a defined direction of a magnitude at least as great as a rigid or semi-rigid support) on the limb or body part that is being used to brace</li> <li>Not used to supply compression therapy (e.g., to reduce size, volume, or swelling of a body member or to help circulation)</li> <li>Not used for convenience or appearance</li> <li>Not used for cosmetic purposes</li> </ul>
Rolling Chair (Geri Chair)		DME	Covered if member meets Mobility Assistive Equipment clinical criteria.  Coverage is limited to those roll-about chairs having casters of at least 5 inches in diameter and officially designed to meet the needs of ill, injured, or otherwise impaired individuals.  Not covered for the wide range of chairs with smaller casters as are found in general use in homes, offices, and institutions for many purposes not related to the care/treatment of ill/injured persons. This type is not primarily medical in nature.  Refer to the:  NCD for Mobility Assistive Equipment (MAE) (280.3)  NCD for Durable Medical Equipment Reference List (280.1)
Safety Rollers			See Walkers.
Sauna Baths		Not Covered	Not primarily medical in nature
Scoliosis Orthosis	Body Jacket Thoracic-lumbar- sacral (TLSO) Cervical-thoracic- lumbar-sacral (CTLSO)	Corrective Appliance/ Orthotic	Coverage criteria apply.  Also see Spinal Orthosis.  General indications for orthotic treatment in idiopathic scoliosis are as follows:  Skeletally immature member's prior to Risser grade 5 (usually one year post menarche in girls)  Children presenting with curvature of 20 to 30 degrees should also be observed, at least initially.  During the observation period, roentgenograms should be obtained at 3 to 6 month intervals and compared with the original films. If the curvature increases by more than 5 degrees in a skeletally immature member, bracing is recommended  Children presenting with 25 degrees to 39 degrees curvature require prompt treatment. These members are at high risk of progression of curvature  Boys with progressive curvature in excess of 25 degrees, including those presenting at Risser grade 4

	ltem		Comments
			<ul> <li>Member's with Scheuermann's Kyphosis including kyphosis of more than 50 degrees. To maintain correction, the brace should be worn until there is improvement in vertebral wedging to roughly 5 degrees. Bracing for longer than 18 months may be necessary to achieve this improvement</li> </ul>
			<b>Note</b> : In very young member bracing may retard progression long enough to allow further trunk growth before the inevitable fusion. Once curvature exceeds 40 degrees, surgical treatment may be the only means of controlling and correcting the deformity.
			Immediate bracing is recommended for the following to allow significant trunk growth prior to surgical intervention:
			<ul> <li>Skeletally immature member's (at Risser grades 0 to 2) presenting with 30 to 40 degrees curvature</li> <li>Flaccid paralysis and 20 degrees or more of curvature</li> </ul>
			<b>Note</b> : Risser grades are based on the degree of bony fusion of the iliac apophysis, from grade 0 (no ossification) to grade 5 (complete bony fusion) (Reamy and Slakey, 2001).
			Note: The recommended duration of bracing varies from 16 hours/day to 23 hours/day.
			Reference Article:
			Schiller JR, Thakur NA, Eberson CP. Brace management in adolescent idiopathic scoliosis. Clin Orthop Relat Res. 2010;468(3):670-678.
			Scoliosis Research Society: Medical Necessity criteria for Scoliosis Bracing (Milwaukee brace; Wilmington brace; Boston brace; Dynamic Spine-Cor brace; Charleston brace; and Providence brace). http://www.srs.org
Shower/Bath Tub	Seat	Not Covered	Not primarily medical in nature
Shower/Bath Tub	) Walk-In	Not Covered	Not primarily medical in nature
Shoes	Inserts/Orthotics	Corrective Appliance/ Orthotic	Refer to the BIP titled Shoes and Foot Orthotics.
	Orthopedic	Corrective Appliance/ Orthotic	
	Prosthetic	Prosthetic	
	Therapeutic (e.g., diabetic shoes)	Corrective Appliance/ Orthotic	

	Item	Coverage	Comments
Sitz Bath (portable	e)	DME	Covered if member has an infection or injury of the perineal area and the item has been prescribed by the patient's physician as a part of his planned regimen of treatment in the patient's home.
Sleep Apnea devi	ce		See Mandibular Device.
Slings		Medical Supply*	Used to support and limit motion of an injured upper arm.
Spinal Orthosis	Cervical-thoracic- lumbar-sacral (CTLSO)	Corrective Appliance/ Orthotic	<ul> <li>Covered when ordered by physician:</li> <li>To reduce pain by restricting mobility of the trunk; or</li> <li>To facilitate healing following an injury to the spine or related soft tissues; or</li> </ul>
	Lumbar-sacral (LSO)		<ul> <li>To facilitate healing following a surgical procedure on the spine or related soft tissue; or</li> <li>To otherwise support weak spinal muscles and/or a deformed spine.</li> </ul>
	Thoracic-lumbar- sacral (TLSO)		Also see <u>Scoliosis Orthosis</u> .
Splints	Bi-directional static progressive stretch splinting (e.g., JAS splints, ERMI system)		Refer to the MMG titled Mechanical Stretching Devices.
	Dental Splint (prefabricated, off- the-shelf bite guard; aka night guard appliance)	Not Covered	Dental splint is an off-the-shelf intraoral device that does not require professional fitting or adjustment and is used to prevent damage to teeth caused by bruxism.  Note: Dental splint does not include oral splints for the treatment of TMJ that require custom fitting and adjustment by a licensed healthcare professional.
	Dynamic (e.g., Dyna Splint)	DME	Refer to the MMG titled Mechanical Stretching Devices.
	Foot (e.g., Dennis- Browne)	Corrective Appliance/ Orthotic	Used as splint/brace to correct rotational anomalies of lower legs; worn during sleep
	Occlusal Splint (custom fabricated bite plate for TMJ)	Corrective Appliance/ Orthotic	Custom made occlusal splints are removable intraoral appliances fabricated and fitted by a licensed healthcare professional to be worn at night for the treatment of painful temporomandibular joint disease.  Refer to the MMG titled Temporomandibular Joint Disorders.  For Sleep Apnea device, see Mandibular Device.
	Wrist/Hand/Finger	Corrective Appliance/ Orthotic	For mild sprains, strains and carpal tunnel conditions; Custom molded covered only when member cannot be fitted with the prefabricated wrist/hand/finger/splint/brace.

l l	ltem		Comments
Standing Frames/Mobile Stander		DME	Covered if Medically Necessary.
			Refer to the InterQual® CP: Durable Medical Equipment, Standing Frames. Click <a href="https://example.com/here">here</a> to view the InterQual® criteria.
			Also refer to the CMS NCD for Mobility Assistive Equipment (MAE) (280.3) and the CMS NCD for Durable Medical Equipment Reference List (280.1).
Stockings	Gradient Compression	Prosthetic	Covered when medical criteria are met; Refer to the InterQual® Medicare: Durable Medical Equipment, Surgical Dressings. Click <a href="https://example.com/here">here</a> to view the InterQual® criteria.
	Stockings (e.g., Jobst stockings)		<b>Note</b> : Coverage is limited to initial 2 pairs of hosiery and replacement of 2 pairs every six month if member is compliant in wearing the hosiery.
	Support Hose (e.g., Ted Hose)	Not Covered	Non-reusable, non-rental item
	Surgical Stockings	Not Covered	Non-reusable, non-rental item
Stump Socks			See Artificial Limb.
Suction Pump or Machine		DME	Covered for members who have difficulty raising and clearing secretions secondary to one of the following:  Cancer or surgery of the throat or mouth  Dysfunction of the swallowing muscles  Unconsciousness or obtunded state
			Tracheostomy Must be appropriate for use without professional supervision
Surgical Boot		Corrective Appliance/ Orthotic	Also known as ambulatory boot.
Syringes	Bulb, Ear	Not Covered	Non re-usable, not rental item
Hypodermic		Medical Supply*	
Telephone Alert Sy	vstem	Not Covered	Not primarily medical in nature
Telephone Arms/Cradle		Not Covered	Not primarily medical in nature
TMJ device			See Occlusal Splint.
ThAIRapy® Vest System			See High Frequency Chest Wall Compression Devices.
Thoracic Lumbar Sacral Orthoses (TLSO)			See <u>Spinal Orthosis</u> ; also see <u>Scoliosis Orthosis</u> .
Tinnitus Masker		Not Covered	Effectiveness not adequately proven.
Toe Filler		Prosthetic	Refer to the BIP titled Shoes and Foot Orthotics.

	ltem		Comments
Toilet Seat, Elevat	Toilet Seat, Elevated Bidet		Not primarily medical in nature, not medical equipment
Tracheostomy	Speaking Valve and Tubes	Prosthetic	A trachea tube has been determined to satisfy the definition of a prosthetic device, and the tracheostomy speaking valve is an add on to the trachea tube which may be considered a medically necessary accessory that enhances the function of the tube, which makes the system a better prosthesis. As such, a tracheostomy speaking valve is covered as an element of the trachea tube which makes the tube more effective.
	Care Kit (Initial and Replacements)	Prosthetic	A tracheostomy care kit is covered for a member following an open surgical tracheostomy which has been open or is expected to remain open for at least 3 months. Replacement kits are covered at one per day only.
Traction Equipment	Cervical (Over-the- Door or Cervical Portable Traction Unit)	DME	<ul> <li>Covered if both the following criteria are met:</li> <li>The member has a musculoskeletal or neurologic impairment requiring traction equipment; and,</li> <li>The appropriate use of a home cervical traction device has been demonstrated to the member and the member tolerated the selected device.</li> </ul>
	Cervical, attached to headboard	Not Covered	No proven clinical advantage compared to over-the-door traction mechanism
	Cervical, not requiring additional stand or frame (e.g., Orthotrac Pneumatic Vest or Pronex)	Not Covered	No proven clinical advantage compared to over-the-door traction mechanism
	Freestanding Traction Stand	Not Covered	No proven clinical advantage compared to over-the-door traction mechanism
	Pneumatic Free- Standing Cervical, Stand/Frame (e.g., Saunders HomeTrac)	DME	Covered if member meets criteria for over-the-door traction unit and one of the following 3 criterion are met:  1. The treating physician orders greater than 20 pounds of cervical traction in the home setting; or, 2. The member has: 3. A diagnosis of temporomandibular joint (TMJ) dysfunction; and 4. Received treatment for the TMJ condition; or, 5. The member has distortion of the lower jaw or neck anatomy (e.g., radical neck dissection) such that a chin halter is unable to be utilized.  Refer to the MMG titled Home Traction Therapy.

ltem		Coverage	Comments
	Power traction equipment/ devices (e.g., VAX-D®, DRX9000, AccuSpineMED™ Spina System™, Lordex® Decompression Unit, DRS System™)		Refer to the MMG titled Motorized Spinal Traction.
	Spinal Unloading Devices (includes spinal and axial decompression units, pneumatic vests)		Refer to the MMG titled Motorized Spinal Traction.
	Weights, bags	DME	When used in conjunction with covered traction services
Transfer (Sliding) Boa	ard	Not Covered	Not primarily medical in nature
Trapeze Bar		DME	A trapeze bar attached to a bed is covered if the member has a covered hospital bed and the member needs this device to sit up because of a respiratory condition, to change body position for other medical reasons, or to get in or out of bed. Not Covered when used on an ordinary bed.  A "Free standing" trapeze equipment is covered if the member does not have a covered hospital bed but the member needs this device to sit up because of a respiratory condition, to change body position for other medical reasons, or to get in or out of bed.
Treadmill Exerciser		Not Covered	Exercise equipment, not primarily medical in nature
Truss		Prosthetic	Covered as prosthetic when used as a holder for surgical dressings or for lumbar strains, sprains or hernia.
Ultraviolet Cabinet		DME	Covered for selected member's with generalized intractable psoriasis.
Unna Boot/Strapping		Medical Supply*	Generally used to treat chronic ulcers that are usually caused by varicosities of the leg.
Urinal (autoclavable)		DME	If member is confined to bed
	Vacuum Assisted Closure Device (VAC) or Negative Pressure Wound Therapy Pump		Refer to the MMG titled Negative Pressure Wound Therapy.
Vacuum Pump or Device (e.g., ErecAid)		Not Covered	Some members may have coverage as DME. Refer to the State Market Plan Enhancements section of the BIP titled Sexual Dysfunction.

ltem		Coverage	Comments
Vaporizers		Not Covered	Not primarily medical in nature
Ventilators (including supplies)		DME	Ventilators (respirators) are recommended for the treatment of neuromuscular diseases, thoracic restrictive diseases, and chronic respiratory failure consequent to chronic obstructive pulmonary disease. This recommendation includes both positive and negative pressure types.  If request is related to Obstructive Sleep Apnea. See <a href="#">CPAP</a> section of DME Grid.
Vitrectomy Face S	Support	DME	Covered following vitrectomy surgery for member who are required to maintain a face down position in the post-operative period.
Walkers (standard)	Rigid (pick-up), adjustable or fixed height	DME	Covered when the following criteria are met:  The member has a mobility limitation that significantly impairs the member's ability to participate in one or more mobility-related activities of daily living (MRADL) in the home.
Folding (pick-up adjustable or fix height Rigid, wheeled, without seat	Folding (pick-up), adjustable or fixed height		A mobility limitation is one that:  • Prevents the member from accomplishing the MRADL entirely, or  • Places the member at recentably determined beightened risk of merbidity or merbidity as merbidity as merbidity.
	_		<ul> <li>Places the member at reasonably determined heightened risk of morbidity or mortality secondary to the attempts to perform the MRADL, or</li> <li>Prevents the member from completing the MRADL within a reasonable time frame; and         <ul> <li>The member is able to safely use the walker; and</li> <li>The functional mobility deficit can be sufficiently resolved with the use of a walker.</li> </ul> </li> <li>*Baskets are not covered</li> </ul>
	Folding, wheeled, without seat		
			Refer to the:  1. DME MAC LCD for Walkers (L33791)  2. NCD for Mobility Assistive Equipment (MAE) (280.3) (Accessed March 18, 2022)  Also refer to the Coverage Summary titled Mobility Assistive Equipment (MAE)
Walkers (special types)	Heavy duty, multiple braking system, variable wheel resistance (Safety Rollers)	DME	Covered for members who meet coverage criteria for a standard walker and who are unable to use a standard walker due to a severe neurologic disorder or other condition causing the restricted use of one hand.  Refer to the:  1. DME MAC LCD for Walkers (L33791) 2. NCD for Mobility Assistive Equipment (MAE) (280.3) (Accessed March 18, 2022) Also refer to the Coverage Summary titled Mobility Assistive Equipment (MAE)
	Heavy duty		Covered for members who meet the coverage criteria for a standard walker and who weigh more than 300 pounds.

ltem		Coverage	Comments
	Leg extensions		Covered only for members 6 feet tall or more.
	With seat		If medically necessary
	With basket	Not Covered	Additional accessories to DME, corrective appliances or prosthetics which are primarily for the comfort or convenience of the member are not covered.
Wearable Cardioverter Defibrillators	Automatic external defibrillator, with integrated electrocardiogram analysis, garment type  Replacement garment for use with automated external defibrillator, each	DME	Covered as DME.
	Replacement electrodes for use with automated external defibrillator, each		
	Replacement battery for automated external defibrillator, each	Not Covered	
Wedge Pillow		Not Covered	Not primarily medical in nature
Wheelchairs and Accessories		DME	Recommended if member meets Mobility Assistive Equipment clinical criteria.  Refer to CMS NCD for Mobility Assistive Equipment (MAE) (280.3).  The following are covered when criteria are met. See the CMS Mobility Assistive Equipment Clinical Criteria for coverage criteria. Wheelchairs are covered only if the member has a DME benefit.  Standard Wheelchair  Lightweight Wheelchair  Specially sized Wheelchair  High Strength Lightweight Wheelchair  High Strength Lightweight Wheelchair  Power Mobility Devices (PMD) which include Power operated vehicle (POV), or scooters, and Power motorized wheelchairs

Item	Coverage	Comments
		<ul> <li>Repairs, replacements and maintenance criteria-refer to the Benefit Interpretation Policy titled         <u>Durable Medical Equipment (DME), Prosthetics, Corrective Appliances/Orthotics (Non-Foot Orthotics) and Medical Supplies.</u></li> <li>Accessories and Options: See the <u>Medicare Local Coverage Determinations (LCD)</u> for Wheelchair Options/Accessories and for Wheelchair Seating.</li> </ul>
Whirlpool Bath Equipment (standard)	DME	<ul> <li>Medical necessity is determined by the following:</li> <li>Evidence that a whirlpool bath offers substantial therapeutic benefit for the member's medical condition</li> <li>Verification that the member is homebound or that treatment in the home is the least costly alternative</li> </ul>
Whirlpool Pump (portable)	Not Covered	Not primarily medical in nature
Wig/Hairpiece (cranial prosthesis)	Not Covered	Not primarily medical in nature  Note: Some Members may have coverage for wigs under the DME benefit Refer to the EOC/ SOB for coverage eligibility.
Wrist splint		See <u>Splints</u> .

<sup>\*</sup>Medical Supplies are covered only when they are incident to a physician's professional services or authorized home health services and are furnished as an integral, although incidental, part of those services in the course of diagnosis or treatment of an injury or illness.

## **Policy History/Revision Information**

Date	State(s) Affected	Summary of Changes
05/01/2023	All	Medical Supplies Grid
		Revised language pertaining to medical necessity clinical coverage criteria; added reference to the:
		Artificial Limbs – Lower Limb, Standard
		<ul> <li>InterQual® Client Defined CP: Durable Medical Equipment, Prosthetic Devices, Wigs, Specialized,</li> <li>Microprocessor or Myoelectric Upper Limbs (Custom) – UHG</li> </ul>
		Oxygen Equipment and Necessary Accessories: Stationary and Portable (Regulated) (e.g.,
		Oxylite, includes Conserver and Tank)
		<ul> <li>InterQual® CP: Durable Medical Equipment, Home Oxygen Therapy</li> </ul>
		Pumps, Infusion and Pain Control
		o InterQual® Medicare: Durable Medical Equipment, External Infusion Pumps
		Standing Frames/Mobile Stander
		o InterQual® CP: Durable Medical Equipment, Standing Frames
		Gradient Compression Stockings (e.g. Jobst stockings)
		<ul> <li>InterQual® Medicare: Durable Medical Equipment, Surgical Dressings</li> </ul>

Date	State(s) Affected	Summary of Changes
		Supporting Information
		Archived previous policy version BIP051.Y

#### **Instructions for Use**

Covered benefits are listed in three (3) sections: Federal/State Mandated Regulations, State Market Plan Enhancements, and Covered Benefits. All services must be medically necessary. Each benefit plan contains its own specific provisions for coverage, limitations, and exclusions as stated in the member's Evidence of Coverage (EOC)/Schedule of Benefits (SOB). If there is a discrepancy between this policy and the member's EOC/SOB, the member's EOC/SOB provision will govern.