Humana

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Medical Coverage Policy

Table of Contents

Related Medical/Pharmacy Coverage Policies Coverage Determination Coding Information Change Summary Description Coverage Limitations <u>References</u>

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Related Medical/Pharmacy Coverage Policies

<u>Code Compendium (Miscellaneous)</u> (bioelectrical impedance analysis [BIA] for body composition) <u>Pneumatic Compression Pumps</u> Support Garments, Fabric

Description

Lymphedema is swelling caused by an abnormal collection of fluid beneath the skin resulting from lymph vessel impairment or lymph node removal. It is generally categorized as primary or secondary; primary is caused by problems with the development of lymph vessels, while secondary is related to something that has damaged the lymph nodes or vessels (eg, surgery, radiation, cancer, infection). Lymphedema differs from edema which is swelling caused by excess fluid that becomes trapped in the body's tissues as a result of medication, pregnancy or underlying disease (eg, heart failure, venous insufficiency, kidney disease, cirrhosis of the liver).

Diagnosis

Assessment and monitoring of lymphedema can be accomplished by a number of methods. One of the most common is circumferential measurement of limb volume. The volume is calculated with measurements obtained with a tape measure at various locations on the limb; it may be compared to

Page: 2 of 13

measurements of the opposite limb. Another method is the water displacement measurement. The limb is submerged into a container of water and the amount that is displaced is measured.

Bioimpedance spectroscopy (BIS), also referred to as bioelectrical impedance analysis, has been proposed as an alternative method to diagnose and/or monitor lymphedema. This device measures the impedance (resistance) of electrical current through extracellular fluid via electrodes that have been attached to the wrist when testing the arm or the ankle when testing the leg. A mild electrical current is passed through the electrode and a measurement of the resistance of the current flow through the fluid is obtained. An example of this device is the **Sozo Digital Health Platform (L-Dex)**. (Refer to Coverage Limitations section)

Treatment

Treatment of lymphedema may be undertaken by a number of methods, either alone or in combination, including, but not limited to, the use of lymphedema garments, manual lymph drainage massage, lymphedema pumps and/or surgery.

Lymphedema garments (also referred to as compression garments), which include sleeves, gloves and stockings, are special bandages that can be worn on the arms, legs, hands or feet to help reduce swelling that is caused by the removal or injury of nearby lymphatic vessels or nodes. The garments provide specific amounts of pressure to keep the fluid from accumulating in the limb. (For more specific information about over-the-counter [OTC] or ready-made garments, please refer to <u>Coverage Limitations section</u>.

Manual lymph drainage massage (also known as complex decongestive physiotherapy or complete decongestive physiotherapy) may be performed by a physical therapist or occupational therapist certified in manual lymph drainage. This technique combines massage, bandaging, exercise and skin care in an attempt to reduce the accumulation of fluid.

Lymphedema pumps (pneumatic compression pumps) are devices that use compressed air to apply pressure to a limb in order to move excess lymph fluid into the rest of the body. A unicompartmental (nonsegmented) device consists of a rubberized sleeve or boot with a single inflatable chamber that exerts uniform pressure along the affected limb. A multicompartmental (segmented) device has multiple chambers in the rubberized sleeve or boot that inflate and deflate in a sequential fashion. These devices may be controlled either with or without manual control of the amount of pressure used in the compartments (manual control is also known as gradient pressure).

An advanced **multicompartmental programmable pneumatic compression device** (formerly referred to as a two-stage multichamber programmable pneumatic compression device) operates similar to the principles of manual lymph drainage (treat the proximal areas first, which is theorized to prepare the distal areas for drainage). Examples of this type of pump include, but may not be limited to, the **AIROS 6, AIROS 8**, **Flexitouch (Flexitouch Plus)** or **Lympha Press Optimal (Lympha Press Optimal Plus)**.

A variation of the multicompartmental pneumatic compression pump is the **CircuFlow 5200 Sequential Compression Device**, which combines intermittent pneumatic compression with a sustained gradient pressure. (**Refer to Coverage Limitations section**)

A new device has been proposed as an alternative treatment for lymphedema, the **Dayspring Active Nonpneumatic Compression System**, which unlike pneumatic compression pumps, does not use air to

Page: 3 of 13

produce the compression, but rather uses a nickel-titanium shape-memory alloy to apply sequential gradient compression. The device is wearable (portable), programmable and battery powered, consisting of the controller and a garment (limb sleeve). It may also be referred to as a nonpneumatic compression device (NPCD). (Refer to Coverage Limitations section)

This policy ONLY addresses treatment for <u>lymphedema</u>. For information regarding **other uses of pneumatic compression pumps** (eg, chronic venous insufficiency, deep vein thrombosis [DVT] prevention), please refer to <u>Pneumatic Compression Pumps</u> Medical Coverage Policy.

Surgery, though not curative and rarely performed, has been suggested as a treatment for those with refractory lymphedema who have not improved with conservative management. Lymphedema surgery may be classified as reconstructive or excisional. Excisional surgical procedures for lymphedema include, but may not be limited to, debulking and liposuction. Reconstructive surgical procedures include, but may not be limited to, microsurgical treatment (eg, microsurgical lymphatico-venous anastomosis, lymphatic-capsular-venous anastomosis, lymphovenous bypass), lymph node transfer (also known as vascularized lymph node transfer) and tissue transfers (eg, omental flap). **(Refer to Coverage Limitations section)**

Coverage Determination

Compression (Lymphedema) Garments

Humana members may be eligible under the Plan for <u>custom-made</u> compression (lymphedema) garments for the extremities (eg, gloves, sleeves or stockings) for the treatment of primary or secondary lymphedema.

Two sets of lymphedema garments per affected extremity are allowed initially; 1 set per affected extremity may be covered thereafter in a <u>rolling 12 month period</u>.* (Sleeves and gloves are separate items; as such, if both should be required for treatment, 2 gloves and 2 sleeves would be allowed initially, with 1 additional of each in subsequent years, if needed.)

*A rolling 12 month period is 12 months after an event, regardless of what month the initial event took place; eg, the initial sets of garments are provided on June 1, 2024, the rolling 12 month period would end on May 31, 2025; in this example, no additional garments would be authorized until June 1, 2025.

Manual Lymph Drainage Massage

Humana members may be eligible under the Plan for **manual lymph drainage massage** (also known as **complex or complete decongestive physiotherapy**) for the treatment of primary or secondary lymphedema when the following criteria are met:

- Individual has undergone a 4 week trial of conservative treatment, including compression garments, elevation of the affected limb and home exercises; **AND**
- The treating healthcare provider determines there has been no improvement or symptoms remain; AND

Page: 4 of 13

- Submission of clinical records documenting the individual's adherence to the conservative treatment that was tried and failed; **AND**
- Treatment is performed by a physical therapist or occupational therapist, preferably certified in manual lymph drainage

This treatment may be applied toward the number of allowable visits of the physical therapy benefit. Refer to specific certificate language regarding physical medicine and rehabilitation services. Most certificates limit the duration or number of visits.

Lymphedema Pumps (Pneumatic Compression Pumps)

Humana members may be eligible under the Plan for the **following types of pneumatic compression pumps** for the treatment of primary or secondary lymphedema of the extremities (arms or legs) when the following criteria are met:

- Initial approval, if criteria are met, is limited to a maximum of 90 days; AND
- Unicompartmental (nonsegmented) or multicompartmental (segmented) lymphedema pump <u>WITHOUT</u> gradient pressure (manual control of the pressure in the chamber) **(E0650, E0651)** of the pressure in the chamber for home use for the treatment of lymphedema when the following are met:
 - Individual has undergone a 4 week trial of conservative therapy, including the use of an appropriate compression garment, exercise and elevation; AND
 - The treating healthcare provider determines there has been no improvement or symptoms remain;
 AND
 - Submission of clinical records documenting the individual's adherence to the conservative therapy that was tried and failed; OR
- Unicompartmental (nonsegmented) or multicompartmental (segmented) lymphedema pump <u>WITH</u> <u>gradient pressure</u> (manual control of the pressure) (E0652)** (manual control of the pressure) when ALL of the following criteria are met:
 - Individual has undergone a 4 week trial of a unicompartmental or multicompartmental lymphedema pump <u>without</u> manual control of the pressure in each chamber; AND
 - $\circ~$ Lymphedema extends from the extremities onto the chest, abdomen or trunk; AND
 - The treating healthcare provider determines that there has been no improvement or symptoms remain; AND

Page: 5 of 13

 Submission of clinical documentation of compliance and adherence with use of the unicompartmental or multicompartmental pump without control of the pressure in each chamber as per the healthcare provider's instructions/prescription

**This includes the advanced multicompartmental programmable pumps (eg, AIROS 6, AIROS 8, Flexitouch [Flexitouch Plus] or Lympha Press Optimal [Lympha Press Optimal Plus]) which are considered equally effective to standard segmented pneumatic compression pumps.

Continuation of Coverage

Lymphedema pumps are initially authorized for 90 days. Continued authorization is dependent upon clinical documentation, submitted by the prescribing healthcare provider, which demonstrates the following:

- Adherence with the use of the device as per the healthcare provider's instructions/prescription; AND
- Confirmation of clinical benefit (eg, improvement in, or prevention in worsening of, the condition for which the device was prescribed)

Coverage Limitations

Humana members may **NOT** be eligible under the Plan for **compression (lymphedema) garments, manual lymph drainage massage or lymphedema pumps** for any indications other than those listed above. All other indications are considered not medically necessary as defined in the member's individual certificate. Please refer to the member's individual certificate for the specific definition.

Humana members may **NOT** be eligible under the Plan for **any other treatment method not listed above** including, but may not be limited to:

- Compression garments for the chest, head, neck or trunk; OR
- Immediate lymphatic reconstruction surgery for prevention of breast cancer-related lymphedema; OR
- Lymphedema pump (and the associated appliance) for treatment of lymphedema *isolated* to the chest or trunk; **OR**
- Lymphedema pump (and the associated appliance) for treatment of lymphedema to the head or neck; **OR**
- Nonpneumatic compression devices (NPCDs) controller (with or without sequential calibrated gradient pressure) or garments including, but not limited to, the **Dayspring nonpneumatic active compression treatment system**; **OR**
- Surgical treatment of lymphedema including, but may not be limited to:

Page: 6 of 13

- Excisional procedures (eg, debulking, liposuction); OR
- Lymph node transfer (also known as vascularized lymph node transfer); OR
- Microsurgical treatment (eg, lymphatico-venous anastomosis, lymphatic-capsular-venous anastomosis, lymphovenous bypass); OR
- Tissue transfer (eg, omental flap)

These are considered experimental/investigational as they are not identified as widely used and generally accepted for the proposed uses as reported in nationally recognized peer-reviewed medical literature published in the English language.

Humana members may **NOT** be eligible under the Plan for **pumps/devices with a sustained gradient pressure while also delivering a higher intermittent pneumatic compression** including, but not limited to, the **CircuFlow 5200 Sequential Compression Device** for any indication. These are considered not medically necessary as defined in the member's individual certificate. Please refer to the member's individual certificate for the specific definition.

Humana members may **NOT** be eligible under the Plan for the use of **bioimpedance spectroscopy** including, but not limited to, the **Sozo (L-Dex)** for diagnosing, monitoring or pre- or postoperative assessment of lymphedema. This is considered experimental/investigational as it is not identified as widely used and generally accepted for the proposed use as reported in nationally recognized peer-reviewed medical literature published in the English language.

Humana members may **NOT** be eligible under the Plan for **ready-made (prefabricated) compression garments/stockings** for any indication. Although they may be prescribed by a health care practitioner, **ready-made compression garments/stockings** are also available without a prescription and may be obtained over-the-counter (OTC) and are therefore generally excluded in the certificate. In the absence of a certificate exclusion for OTC items, **ready-made compression garments/stockings** are considered not medically necessary as defined in the member's individual certificate. Please refer to the member's individual certificate for the specific definition.

Coding Information

Any codes listed on this policy are for informational purposes only. Do not rely on the accuracy and inclusion of specific codes. Inclusion of a code does not guarantee coverage and/or reimbursement for a service or procedure.

Page: 7 of 13

| CPT® Code(s) | Description | Comments |
|---|--|---|
| 38308 | Lymphangiotomy or other operations on lymphatic channels | Not Covered if used to report surgical treatment of lymphedema |
| 93702 | Bioimpedance spectroscopy (BIS), extracellular fluid analysis for lymphedema assessment(s) | Not Covered |
| 97016 | Application of a modality to 1 or more areas; vasopneumatic devices | |
| 97140 | Manual therapy techniques (eg, mobilization/ manipulation, manual lymphatic drainage, manual traction), 1 or more regions, each 15 minutes | |
| CPT [®] Category III Code(s) | Description | Comments |
| No code(s) id | lentified | |
| HCPCS Code(s) | Description | Comments |
| A6549 | Gradient compression stocking/sleeve, not otherwise specified | Not Covered if used to report ready-made (prefabricated) compression garments/stockings (ie, Over the counter) |
| A6567 | Gradient compression garment, neck/head, custom, each | Not Covered New Code Effective 01/01/2024 |
| E0650 | Pneumatic compressor, nonsegmental home model | |
| E0651 | Pneumatic compressor, segmental home model without calibrated gradient pressure | |
| E0652 | Pneumatic compressor, segmental home model with calibrated gradient pressure | Not Covered if used to report any pump/device outlined in Coverage Limitations section |
| E0655 | Nonsegmental pneumatic appliance for use with pneumatic compressor, half arm | |
| E0656 | Segmental pneumatic appliance for use with pneumatic compressor, trunk | Not Covered |
| E0657 | Segmental pneumatic appliance for use with pneumatic compressor, chest | Not Covered |

| E0660 | Nonsegmental pneumatic appliance for use with pneumatic compressor, full leg | |
|-------|--|---|
| E0665 | Nonsegmental pneumatic appliance for use with pneumatic compressor, full arm | |
| E0666 | Nonsegmental pneumatic appliance for use with pneumatic compressor, half leg | |
| E0667 | Segmental pneumatic appliance for use with pneumatic compressor, full leg | Not Covered if used to report any pump/device outlined in Coverage Limitations section |
| E0668 | Segmental pneumatic appliance for use with pneumatic compressor, full arm | Not Covered if used to report any pump/device outlined in Coverage Limitations section |
| E0669 | Segmental pneumatic appliance for use with pneumatic compressor, half leg | Not Covered if used to report any pump/device outlined in Coverage Limitations section |
| E0670 | Segmental pneumatic appliance for use with pneumatic compressor, integrated, two full legs and trunk | Not Covered |
| E0671 | Segmental gradient pressure pneumatic appliance, full leg | Not Covered if used to report any pump/device outlined in Coverage Limitations section |
| E0672 | Segmental gradient pressure pneumatic appliance, full arm | Not Covered if used to report any pump/device outlined in Coverage Limitations section |
| E0673 | Segmental gradient pressure pneumatic appliance, half leg | Not Covered if used to report any pump/device outlined in Coverage Limitations section |
| E0676 | Intermittent limb compression device (includes all accessories), not otherwise specified | |
| E0677 | Non-pneumatic sequential compression garment, trunk | Not Covered |
| | Non-pneumatic sequential compression garment, full leg | Not Covered |
| E0678 | | New Code Effective 01/01/2024 |

Page: 9 of 13

| | | Not Covered |
|-------|---|--|
| E0679 | Non-pneumatic sequential compression garment, half leg | New Code Effective 01/01/2024 |
| E0680 | Non-pneumatic compression controller with sequential calibrated gradient pressure | Not Covered New Code Effective 01/01/2024 |
| E0681 | Non-pneumatic compression controller without calibrated gradient pressure | Not Covered New Code Effective 01/01/2024 |
| E0682 | Non-pneumatic sequential compression garment, full arm | Not Covered New Code Effective 01/01/2024 |
| E1399 | Durable medical equipment, miscellaneous | Not Covered if used to report any treatments outlined in Coverage Limitations section |
| L8010 | Breast prosthesis, mastectomy sleeve | |
| S8420 | Gradient pressure aid (sleeve and glove combination), custom made | |
| S8422 | Gradient pressure aid (sleeve), custom made, medium weight | |
| S8423 | Gradient pressure aid (sleeve), custom made, heavy weight | |
| S8425 | Gradient pressure aid (glove), custom made, medium weight | |
| S8426 | Gradient pressure aid (glove), custom made, heavy weight | |
| S8429 | Gradient pressure exterior wrap | |
| S8950 | Complex lymphedema therapy, each 15 minutes | |

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Lymphedema – Diagnosis and Treatment Page: 11 of 13

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Page: 13 of 13

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Change Summary

- 06/06/2024 Annual Review, No Coverage Change.