MEDICAL POLICY

SUBJECT: LIMB PNEUMATIC COMPRESSION DEVICES FOR VENOUS THROMBOEMBOLISM PROPHYLAXIS

POLICY NUMBER: 1.01.51
CATEGORY: Equipment/ Supplies

EFFECTIVE DATE: 06/27/13
REVISED DATE: 06/26/14, 09/15/15, 09/21/17.

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If the member's subscriber contract excludes coverage for a specific service it is not covered under that contract. In such cases, medical policy criteria are not applied.

Medical policies apply to commercial and Medicaid products only when a contract benefit for the specific service exists.

Medical policies only apply to Medicare products when a contract benefit exists and where there are no National or Local Medicare coverage decisions for the specific service.

POLICY STATEMENT:

I. Based on our criteria, pneumatic compression devices have been medically proven to be effective and are considered medically appropriate when used in the home for deep vein thrombosis prophylaxis after major surgery, including major orthopedic surgery, in patients when pharmacological prophylaxis is contraindicated. Use of pneumatic compression device is allowed for up to 30 days post-operatively.

II. Based upon our criteria and assessment of peer-reviewed literature, use of pneumatic compression devices for prevention of deep vein thrombosis, other than as described in Policy Statement I, is considered not medically necessary.

Refer to Corporate Medical Policy 1.01.17 Pneumatic Compression Devices/Lymphedema Pumps

POLICY GUIDELINES:

I. Major orthopedic surgery includes total hip arthroplasty (THA), total knee arthroplasty (TKA), or hip fracture surgery (HFS).

II. American College of Chest Physicians (ACCP) guidelines on prevention of VTE in orthopedic surgery patients list the following general risk factors for bleeding:
   A. Previous major bleeding (and previous bleeding risk similar to current risk),
   B. Severe renal failure
   C. Concomitant antiplatelet agent, and
   D. Surgical factors: history of or difficult-to-control surgical bleeding during the current operative procedure, extensive surgical dissection, and revision surgery.
   E. Thrombocytopenia

III. The Federal Employee Health Benefit Program (FEHBP/FEP) requires that procedures, devices or laboratory tests approved by the U.S. Food and Drug Administration (FDA) may not be considered investigational and thus these procedures, devices or laboratory tests may be assessed only on the basis of their medical necessity.

DESCRIPTION:

Pneumatic compression devices have been utilized in the post-operative period for prevention of deep vein thrombosis. The devices include various types of wraps for the arms or legs and a programmable control module. The wraps of some of these devices are capable of providing cooling or heating to the extremity. Examples of devices which can be used in the home after discharge include the VascuTherm2 (Thermotek, Inc) and the Triple Play VT® (Compression Solutions, Inc).

RATIONALE:

Anticoagulation is the mainstay of DVT prophylaxis after major surgery and is sometimes continued into the outpatient setting. Treatment with pneumatic compression devices may offer an additional benefit when used in conjunction with anticoagulation in the inpatient setting, but is not commonly used in the outpatient setting. Studies involving the use of compression devices post-operatively are limited; the studies are often small, nonrandomized with considerable variation in the comparison studies by type of compression stocking and intermittent compression device used, patient group, DVT detection method, and prophylaxis protocol. Many of the studies are in the setting of the hospital rather than...
outpatient, consequently conclusions from the hospital setting may not be able to be applied to the outpatient setting. This is due to differences in the levels of mobility in the hospital versus the outpatient setting, which may change the risk for DVT. Also, the use of pneumatic compression devices in the hospital can be more highly controlled and monitored. In the outpatient setting, there are questions about the degree of compliance with the devices, including the ability to correctly use them in the absence of professional supervision.

In 2011, the American Academy of Orthopedic Surgeons (AAOS) published an updated guideline on prevention of venous thromboembolism in patients undergoing elective hip and knee arthroplasty. The work group recommended the use of pharmacologic agents and/or mechanical compressive devices for the prevention of venous thromboembolism in patients undergoing elective hip or knee arthroplasty who are not at elevated risk beyond that of the surgery itself for venous thromboembolism or bleeding (consensus recommendation), and who have also had a previous venous thromboembolism ( consensus recommendation). For those patients undergoing elective hip or knee arthroplasty, who also have a known bleeding disorder (e.g., hemophilia) and/or active liver disease, mechanical compressive devices for preventing venous thromboembolism is recommended (consensus recommendation).

The American College of Chest Physicians (ACCP) (2012) recommended the use of limb compression devices in non-orthopedic general and abdominal-pelvic surgical patients, in patients who were classified as low risk for VTE and moderate risk for VTE and high-risk for bleeding rather than no prophylaxis and as an option for those individuals with moderate and high risk for VTE and who were not at high risk for bleeding in addition to pharmacologic prophylaxis.

In 2007, the American College of Obstetricians-Gynecologists (ACOG) published a practice bulletin on prevention of DVT and PE after gynecologic surgery. As with the ACCP recommendations, described above, prophylaxis recommendations varied according to patient risk level. For patients at moderate and high risk of DVT, intermittent pneumatic compression was one of the recommended options for DVT prophylaxis. For patients at highest risk i.e., older than 60 years plus prior VTE, cancer or molecular hypercoagulable state, IPC or graduated compression stockings plus LDUH or LMWH was recommended as a prophylaxis option. The devices should be used continuously until ambulation and discontinued only at the time of hospital discharge except for the highest risk patients, who should consider prophylaxis for 2-4 weeks after discharge.

A Cochrane review by Naccarato, et al. (2010), identified 2 clinical trials and 2 small studies using gradient compression stockings (GCS) or intermittent pneumatic compression (IPC) devices for DVT prophylaxis in stroke patients. The authors concluded that evidence from randomized trials does not support the routine use of GCS to reduce the risk of DVT after acute stroke. There is insufficient evidence to support the routine use of IPC to reduce the risk of DVT in acute stroke and further larger randomized studies of IPC are needed to reliably assess the balance of risks and benefits of this intervention.

**CODES:**

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<th>Number</th>
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<td>Eligibility for reimbursement is based upon the benefits set forth in the member’s subscriber contract.</td>
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CODES MAY NOT BE COVERED UNDER ALL CIRCUMSTANCES. PLEASE READ THE POLICY AND GUIDELINES STATEMENTS CAREFULLY.

Codes may not be all inclusive as the AMA and CMS code updates may occur more frequently than policy updates.

**CPT:** No code(s)

**HCPCS:**

- 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
- E0660 Nonsegmental pneumatic appliance for use with pneumatic compressor, full
E0666  leg  half leg
E0667  Segmental pneumatic appliance for use with pneumatic compressor, full leg half leg
E0669  Segmental gradient pressure pneumatic appliance, full leg half leg
E0671  Pneumatic compression device, high pressure, rapid inflation/deflation cycle for arterial insufficiency (unilateral or bilateral system)
E0673  Intermittent limb compression device (includes all accessories), not otherwise specified

ICD9:
714.0-714.9  Rheumatoid arthritis and other inflammatory polyarthropathies (code range)
715.00-715.98  Osteoarthritis (code range)
716.15- 716.16  Traumatic arthropathy (hip and knee codes respectively)
733.14  Pathologic fracture of neck of femur
820.00-820.9  Fracture of neck of femur (code range)
821.00-821.11  Fracture of shaft of femur (code range)
V43.64-V43.65  Organ or tissue replaced by other means, joint (hip and knee codes respectively)
V54.81  Aftercare following joint replacement

ICD10:
M05.00-M05.09  Felty's syndrome (code range)
M05.20-M05.29  Rheumatoid vasculitis with rheumatoid arthritis (code range)
M05.30-M05.39  Rheumatoid heart disease with rheumatoid arthritis (code range)
M05.40-M05.49  Rheumatoid myopathy with rheumatoid arthritis (code range)
M05.50-M05.579  Rheumatoid polyneuropathy with rheumatoid arthritis (code range)
M05.60-M05.79  Rheumatoid arthritis of unspecified site with or without involvement of other organs and systems (code range)
M05.80-M06.09  Rheumatoid arthritis with or without rheumatoid factor (code range)
M06.1  Adult-onset Still's disease
M062.0-M06.29  Rheumatoid bursitis (code range)
M06.30-M06.39  Rheumatoid nodule (code range)
M06.4  Inflammatory polyarthropathy
M06.80-M06.9  Other specified rheumatoid arthritis (code range)
M08.00-M08.29  Unspecified juvenile rheumatoid arthritis (code range)
M08.3  Juvenile rheumatoid polyarthritis (seronegative)
M08.40-M08.48  Pauciarticular juvenile rheumatoid arthritis (code range)
M08.80-M08.99  Other juvenile arthritis (code range)
REFERENCES:


**KEY WORDS:**
Venodyne, VascuTherm2, Triple Play VT®

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**CMS COVERAGE FOR MEDICARE PRODUCT MEMBERS**

There is currently a National Coverage Determination (NCD) and a Local Coverage Determination (LCD) for Pneumatic Compression Devices. Please refer to the following websites for Medicare Members:

**NCD SITE:**

**LCD SITE:**
http://www.cms.gov/medicare-coverage-database/details/lcd-details.aspx?LCDId=11503&ContrId=137&ver=29&ContrVer=1&CntrctrSelected=137*1&Cntrctr=137&name=NHIC%2c+Corp.+(16003%2c+DME+MAC)&s=41&DocType=All&bc=AggAAAIAAAAAA%3d%3d&