

Negative Pressure Wound Therapy DME-011

Iowa Medicaid Program:	Prior Authorization	Effective Date:	9/11/2009
Revision Number:	9	Last Rev Date:	10/20/2023
Reviewed By:	Medicaid Medical Director	Next Rev Date:	10/18/2024
Approved By:	Medicaid Clinical Advisory Committee	Approved Date:	6/4/2018

Criteria

Negative pressure wound therapy (NPWT), also called vacuum-assisted wound closure, refers to wound dressing systems that continuously or intermittently apply sub-atmospheric pressure to the system, which provides a positive pressure to the surface of a wound. NPWT has become a popular treatment modality for the management of many acute and chronic wounds. This therapy should be considered an adjunct to the basic principles of wound care.

NPWT exerts its effect through direct and indirect effects of sub-atmospheric pressure. These effects include stabilization of the wound environment, increased blood flow, and deformation of the wound. Deformation is a powerful stimulus for cellular processes that stimulate granulation tissue and accelerate wound healing. The dressing and tubing are typically changed every 48 to 120 hours (2 to 5 days) depending upon the clinical situation.

Both systemic and local wound factors can contribute to delayed wound healing. Systemic factors (e.g., poor nutrition, wound ischemia) should be identified and corrected to the extent that is possible. Local wound factors that interfere with normal healing include desiccation, tissue edema, excessive exudate, poor tissue apposition (e.g., grafts and flaps), and wound infection. Stagnant fluid is associated with cytogenetic factors that impede wound healing.

NPWT has been applied to a wide range of clinical situations, including the open abdomen, following surgical debridement of acute or chronic wounds (e.g., orthopedic, necrotizing infection, pressure ulcer), diabetic foot ulcers, and reconstructive surgery (e.g., burns, skin graft, muscle flap). It has also been used to prevent surgical wound infection (prophylaxis) and as a means of instillation therapy.

Contraindications

NPWT should not be used when any of the following situations are present:

- Exposed vital structures NPWT, in the presence of exposed organs, blood vessels, or vascular grafts, increases the risk for tissue erosion, which can lead to enteric fistula formulation or hemorrhage. NPWT is generally avoided until an intervening granulation layer or tissue flap, or graft provides coverage. Although some clinicians report success using barrier dressings, caution is advised when implementing this practice.
- Presence of malignant tissue As with normal tissues, growth of malignant tissue is promoted in the presence of sub-atmospheric pressure. Malignant tissue is also more friable and prone to bleeding.

The use of non-powered (mechanical) NPWT devices (e.g., the Smart Negative Pressure [SNaP] Wound Care System) is considered experimental and investigational because their effectiveness has not been established.

Criteria

NWPT is considered medically necessary when **ALL** the following have been met:

- A complete wound care program, which includes <u>ALL</u> the following has been implemented:
 - a. Documentation of evaluation, care, and wound measurements by a licensed medical professional; **AND**
 - b. Application of dressings to maintain a moist environment; **AND**
 - c. Debridement of necrotic tissue when present; **AND**
 - d. Underlying nutritional deficiencies have been addressed; **AND**
 - e. Underlying comorbidities, such as diabetes or venous insufficiency, have been managed; **AND**
- 2. **ONE** or more of the following eligible conditions is present:
 - a. Stage III or IV pressure ulcers; **OR**
 - b. Neuropathic ulcers; **OR**
 - c. Ulcers related to venous or arterial insufficiency; **OR**
 - d. Wound dehiscence with exposed hardware or bone; **OR**
 - e. Post sternotomy wound infection or mediastinitis; OR
 - f. Complications of a surgically created wound where accelerated granulation therapy is necessary and cannot be achieved by other available topical wound treatment;

 AND
- 3. None of the following **contraindications** to NPWT are present:
 - a. Exposed anastomotic site; **OR**
 - b. Exposed nerve; **OR**
 - c. Exposed organs; OR
 - d. Exposed vasculature; OR
 - e. Presence of cancer in the wound; **OR**

- f. Necrotic tissue with eschar: **OR**
- g. Non-enteric and unexplored fistulas; OR
- h. Untreated osteomyelitis.

NWPT for continued use is considered medically necessary when all of the following are met:

- Assessment of wound dimensions and characteristics are performed by a licensed healthcare professional on at least a monthly basis and documented in the medical record; AND
- 2. The measurements document progressive wound healing.

Definitions – Pressure Ulcer Stages

Pressure Injury:

A pressure injury is localized damage to the skin and/or underlying soft tissue usually over a bony prominence or related to a medical or other device. The injury can present as intact skin or an open ulcer and may be painful. The injury occurs because of intense and/or prolonged pressure or pressure in combination with shear. The tolerance of soft tissue for pressure and shear may also be affected by microclimate, nutrition, perfusion, co-morbidities, and condition of the soft tissue.

Stage I: Intact skin with a localized area of non-blanchable erythema, which may appear differently in darkly pigmented skin. Presence of blanchable erythema or changes in sensation, temperature, or firmness may precede visual changes. Color changes do not include purple or maroon discoloration; these may indicate deep tissue pressure injury.

Stage 2: Partial-thickness loss of skin with exposed dermis. The wound bed is viable, pink or red, moist, and may also present as an intact or ruptured serum-filled blister. Adipose (fat) is not visible and deeper tissues are not visible. Granulation tissue, slough, and eschar are not present. These injuries commonly result from adverse microclimate and shear in the skin over the pelvis and shear in the heel. This stage should not be used to describe moisture associated skin damage including incontinence associated dermatitis, intertriginous dermatitis, medical adhesive related skin injury, or traumatic wounds (skin tears, burns, abrasions).

Stage 3: Full-thickness loss of skin, in which adipose (fat) is visible in the ulcer and granulation tissue with rolled wound edges are often present. Slough and/or eschar may be visible. The depth of tissue damage varies by anatomical location; areas of significant adiposity can develop deep wounds. Undermining and tunneling may occur. Fascia, muscle, tendon, ligament, cartilage and/or bone are not exposed. If slough or eschar obscures the extent of tissue loss this is an unstageable pressure injury.

Stage 4: Full-thickness skin and tissue loss with exposed or directly palpable fascia, muscle, tendon, ligament, cartilage, or bone in the ulcer. Slough and/or eschar may be visible with rolled edges, undermining and/or tunneling often occur. Depth varies by anatomical location. If slough or eschar obscures the extent of tissue loss this is an unstageable pressure injury.

Coding

The following list of codes is provided for reference purposes only and may not be all inclusive. The inclusion of a code does not imply any right to reimbursement or guarantee claim payment, nor does the exclusion of a code imply that its association to the HCPCS/CPT code is inappropriate.

HCPCS	Description
E2402	Negative pressure wound therapy electrical pump, stationary or portable.

References

Optum 360/EncoderPro.

Gestring M. Negative Pressure Wound Therapy. UpToDate. Last Updated January 03, 2022.

Negative Pressure Wound Therapy Pumps – Policy Article. Medicare Coverage Database. Article ID A52511. Revision Effective Date: August 15, 2021.

Article - Negative Pressure Wound Therapy Pumps - Policy Article (A52511) (cms.gov).

Negative Pressure Wound Therapy Pumps. Local Coverage Determination L33821. For services performed on or after May 1, 2021.

LCD - Negative Pressure Wound Therapy Pumps (L33821) (cms.gov).

Hingorani A. LaMuraglia GM. Henke P. et al. The management of diabetic foot: a clinical practice guideline by the Society for Vascular Surgery in collaboration with the American Podiatric Medical Association and the Society for Vascular Medicine. J Vasc Surg. 2016; 63(2 Suppl):3S-21S.

Negative Pressure Wound Therapy. Milliman Clinical Guidelines Ambulatory Care. ACG: A-0346. 25th Edition. Last updated June 7, 2021.

National Pressure Injury Advisory Panel (NPIAD). Prevention and treatment of pressure ulcers: clinical practice guideline. 3rd edition. 2019.

http://www.internationalguideline.com/guideline.

Development of utilization management criteria may also involve research into other state Medicaid programs, other payer policies, consultation with experts and review by the Medicaid Clinical Advisory Committee (CAC). These sources may not be referenced individually unless they are specifically published and are otherwise applicable to the criteria at issue.

Change Date	Changed By	Description of Change	Version
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11/14/2023	Medical Director	MMIS updated.	10
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Change Date	Changed By	Description of Change	Version
10/20/2023	CAC	Annual review. Awaiting MMIS updates.	9
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10/21/2022			_
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4/17/2015	CAC	Added paragraph in References.	3
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Change Date	Changed By	Description of Change	Version
		Changed name from Wound Vacuum to	
		Negative Pressure Wound Therapy (NPWT).	
4/18/2014	Medical Director	Added under Coverage Position (Outpatient)	2
		#1 "all wound measurements must be	
		provided". Formatting changes.	
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Change Date	Changed By	Description of Change	Version
		Removed #1 and replaced as definition of	
		chronic wounds. Re-number #2 and #3 to be	
		#1 and #2. Removed "of the healing arts" from	
10/27/2012	CAC	criterion #2.	1
		Coverage positions added "Patient must meet	
		ONE of the following:". Removed wound/ulcer	
		"with lack of healing" for outpatient setting #1.	