

Medical Coverage Policy | Mechanical Wound Suction



EFFECTIVE DATE: 12|01|2014
POLICY LAST UPDATED: 11|04|2014

OVERVIEW

Disposable non-powered mechanical negative pressure wound therapy or single use non-electrically powered negative pressure wound therapy have been proposed for the treatment of smaller wounds. These devices can be used in the hospital, outpatient and/or home settings.

PRIOR AUTHORIZATION

BlueCHiP for Medicare and Commercial

Prior Authorization review is not required.

POLICY STATEMENT

BlueCHiP for Medicare

Use of nonpowered NPWT systems for the treatment of acute or chronic wounds is **not covered**, as they do not meet the DME benefit durability requirement.

Commercial

Use of nonpowered NPWT systems for the treatment of acute or chronic wounds is considered **not medically necessary**, as there is insufficient peer-reviewed literature to support the efficacy of the service.

MEDICAL CRITERIA

Not applicable.

BACKGROUND

The management and treatment of chronic wounds, including decubitus ulcers, remain a treatment challenge. Most chronic wounds will heal only if the underlying cause, i.e., venous stasis, pressure, infection, etc., is addressed. In addition, cleaning the wound to remove nonviable tissue, microorganisms, and foreign bodies is essential to create the optimal conditions for either re-epithelialization (i.e., healing by secondary intention) or preparation for wound closure with skin grafts or flaps (i.e., healing by primary intention). Therefore, debridement, irrigation, whirlpool treatments, and wet-to-dry dressings are common components of chronic wound care.

A nonpowered (mechanical) NPWT system has been developed; 1 device is the Smart Negative Pressure (SNaP) Wound Care System. This device is portable and lightweight (3 oz) and can be worn underneath clothing. This system consists of a cartridge, dressing, and strap; the cartridge acts as the negative pressure source. The system is reported to generate negative pressure levels similar to other NPWT systems. This system is fully disposable.

A nonpowered NPWT device, the SNaP® Wound Care System from Spiracur, is a Class II device requiring notification to market but not having FDA premarket approval. It received 510(k) marketing clearance from FDA in 2009 (K081406) and is designed to remove small amounts of exudate from chronic, traumatic, dehisced, acute, subacute wounds and diabetic and pressure ulcers.

Reports with small numbers of patients using the non-powered (mechanical) gauze-based NPWT system are insufficient to draw conclusions about its impact on net health outcome, both for the device itself and in comparison with current care. There are important unanswered questions about efficacy and tolerability.

Well-designed comparative studies with larger numbers of patients are needed. Since the impact on net health outcome compared to existing technology is not known, non-powered (mechanical) NPWT is considered not medically necessary as there is no proven efficacy.

COVERAGE

Benefits may vary between groups/contracts. Please refer to the appropriate Evidence of Coverage or Subscriber Agreement for limitations of benefits/coverage when services are not medically necessary.

CODING

The following code is **not medically necessary for Commercial** and **not covered for BlueCHiP for Medicare**, as it does not meet the DME benefit durability requirement:

A9272

The following codes are considered **not covered for BlueCHiP for Medicare**, as they do not meet the DME benefit durability requirement. **They are not applicable to Commercial, as an alternate code should be used.**

G0456, G0457

RELATED POLICIES

Preauthorization via Web-Based Tool for Durable Medical Equipment (DME)

PUBLISHED

Provider Update	Jan	2015
Provider Update	Feb	2014
Provider Update	Feb	2013
Provider Update	Sep	2011
Provider Update	Dec	2010
Provider Update	Feb	2009
Policy Update	Sep	2008

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