

Medical Policy



Surgical Dressings

▼ Description

A Surgical dressing is a cover over a wound or surgical incision. Examples of wounds include but are not limited to: cellulitis, ulcerated wounds, stasis ulcers, or burns.

▼ Policy

If the coverage criteria described below are not met, the claim will be denied as not reasonable and necessary.

▼ Policy Guidelines

Surgical dressings are covered for as long as they are medically necessary. Dressings over a percutaneous catheter or tube (e.g., intravascular, epidural, nephrostomy, etc.) are covered as long as the catheter or tube remains in place and after removal until the wound heals.

Surgical dressings used in conjunction with investigational wound healing therapy (e.g., platelet derived wound healing formula) may be covered if all applicable coverage criteria are met based on the number and type of surgical dressings that are appropriate to treat the wound if the investigational therapy were not being used.

When a wound cover with an adhesive border is being used, no other dressing is needed on top of it and additional tape is usually not required. Reasons for use of additional tape must be well documented. An adhesive border is usually more binding than that obtained with separate taping and is therefore indicated for use with wounds requiring less frequent dressing changes.

Use of more than one type of wound filler or more than one type of wound cover in a single wound is rarely medically necessary and the reasons must be well documented. An exception is an alginate or other fiber gelling dressing wound cover or a saline, water, or hydrogel impregnated gauze dressing which might need an additional wound cover.

It may not be appropriate to use some combinations of a hydrating dressing on the same wound at the same time as an absorptive dressing (e.g., hydrogel and alginate).

Because composite dressings, foam and hydrocolloid wound covers, and transparent film, when used as secondary dressings, are meant to be changed at frequencies less than daily, appropriate clinical judgment should be used to avoid their use with primary dressings which require more frequent dressing changes. When claims are submitted for these dressings for changes greater than once every other day, the quantity in excess of that amount will be denied as not reasonable and necessary. While a highly exudative wound might require such a combination initially, with continued proper management the wound usually progresses to a point where the appropriate selection of these products results in the less frequent dressing changes which they are designed to allow. An example of an inappropriate combination is the use of a specialty absorptive dressing on top of non-impregnated gauze being used as a primary dressing.

Dressing size must be based on and appropriate to the size of the wound. For wound covers, the pad size is usually about 2 inches greater than the dimensions of the wound. For example, a 5 cm x 5 cm (2 in. x 2 in.) wound requires a 4 in. x 4 in. pad size.

The quantity and type of dressings dispensed at any one time must take into account the current status of the wound(s), the likelihood of change, and the recent use of dressings.

Dressing needs may change frequently (e.g., weekly) in the early phases of wound treatment and/or with heavily draining wounds. Suppliers are also expected to have a mechanism for determining the quantity of dressings that the patient is actually using and to adjust their provision of dressings accordingly. No more than a one month's supply of dressings may be provided at one time, unless there is documentation to support the necessity of greater quantities in the home setting in an individual case. An even smaller quantity may be appropriate in the situations described above.

Surgical dressings must be tailored to the specific needs of an individual patient. When surgical dressings are provided in kits, only those components of the kit that meet the definition of a surgical dressing, that are ordered by the physician, and that are medically necessary are covered.

The following are some specific coverage guidelines for individual products when the products themselves are necessary in the individual patient. The medical necessity for more frequent change of dressing must be documented in the patient's medical record and submitted with the claim.

ALGINATE OR OTHER FIBER GELLING DRESSING (A6196-A6199):

Alginate or other fiber gelling dressing covers are covered for moderately to highly exudative full thickness wounds (e.g., stage III or IV ulcers); and alginate or other fiber gelling dressing fillers for moderately to highly exudative full thickness wound cavities (e.g., stage III or IV ulcers). They are not medically necessary on dry wounds or wounds covered with eschar. Usual dressing

change is up to once per day. One wound cover sheet of the approximate size of the wound or up to 2 units of wound filler (1 unit = 6 inches of alginate or other fiber gelling dressing rope) is usually used at each dressing change. It is usually inappropriate to use alginates or other fiber gelling dressings in combination with hydrogels.

COMPOSITE DRESSING (A6203-A6205):

Usual composite dressing change is up to 3 times per week, one wound cover per dressing change.

CONTACT LAYER (A6206-A6208):

Contact layer dressings are used to line the entire wound; they are not intended to be changed with each dressing change. Usual dressing change is up to once per week.

FOAM DRESSING (A6209-A6215):

Foam dressings are covered when used on full thickness wounds (e.g., stage III or IV ulcers) with moderate to heavy exudate. Usual dressing change for a foam wound cover used as a primary dressing is up to 3 times per week. When a foam wound cover is used as a secondary dressing for wounds with very heavy exudate, dressing change may be up to 3 times per week. Usual dressing change for foam wound fillers is up to once per day.

GAUZE, NON-IMPREGNATED (A6216-A6221, A6402-A6404, A6407):

Usual non-impregnated gauze dressing change is up to 3 times per day for a dressing without a border and once per day for a dressing with a border. It is usually not necessary to stack more than 2 gauze pads on top of each other in any one area.

GAUZE, IMPREGNATED, WITH OTHER THAN WATER, NORMAL SALINE, HYDROGEL, OR ZINC PASTE (A6222-A6224, A6266):

Usual dressing change for gauze dressings impregnated with other than water, normal saline, hydrogel, or zinc paste is up to once per day.

GAUZE, IMPREGNATED, WATER OR NORMAL SALINE (A6228-A6230):

There is no medical necessity for these dressings compared to non-impregnated gauze which is moistened with bulk saline or sterile water. When these dressings are billed, they will be denied as not reasonable and necessary.

HYDROCOLLOID DRESSING (A6234-A6241):

Hydrocolloid dressings are covered for use on wounds with light to moderate exudate. Usual dressing change for hydrocolloid wound covers or hydrocolloid wound fillers is up to 3 times per week.

HYDROGEL DRESSING (A6231-A6233, A6242-A6248):

Hydrogel dressings are covered when used on full thickness wounds with minimal or no exudate (e.g., stage III or IV ulcers). Hydrogel dressings are not usually medically necessary for stage II ulcers. Documentation must substantiate the medical necessity for use of hydrogel dressings for stage II ulcers (e.g., location of ulcer is sacro-coccygeal area). Usual dressing change for hydrogel wound covers without adhesive border or hydrogel wound fillers is up to once per day. Usual dressing change for hydrogel wound covers with adhesive border is up to 3 times per week.

The quantity of hydrogel filler used for each wound must not exceed the amount needed to line the surface of the wound. Additional amounts used to fill a cavity are not medically necessary. Documentation must substantiate the medical necessity for code A6248 billed in excess of 3 units (fluid ounces) per wound in 30 days.

Use of more than one type of hydrogel dressing (filler, cover, or impregnated gauze) on the same wound at the same time is not medically necessary.

SPECIALTY ABSORPTIVE DRESSING (A6251-A6256):

Specialty absorptive dressings are covered when used for moderately or highly exudative wounds (e.g., stage III or IV ulcers). Usual specialty absorptive dressing change is up to once per day for a dressing without an adhesive border and up to every other day for a dressing with a border.

TRANSPARENT FILM (A6257-A6259):

Transparent film dressings are covered when used on open partial thickness wounds with minimal exudate or closed wounds. Usual dressing change is up to 3 times per week.

WOUND FILLER, NOT ELSEWHERE CLASSIFIED (A6261-A6262):

Usual dressing change is up to once per day.

WOUND POUCH (A6154):

Usual dressing change is up to 3 times per week.

TAPE (A4450,A4452):

Tape is covered when needed to hold on a wound cover, elastic roll gauze or non-elastic roll gauze. Additional tape is usually not required when a wound cover with an adhesive border is used. The medical necessity for tape in these situations must be documented. Tape change is determined by the frequency of change of the wound cover. Quantities of tape submitted must reasonably reflect the size of the wound cover being secured. Usual use for wound covers measuring 16 square inches or less is up to 2 units per dressing change; for

wound covers measuring 16 to 48 square inches, up to 3 units per dressing change; for wound covers measuring greater than 48 square inches, up to 4 units per dressing change.

LIGHT COMPRESSION BANDAGE (A6448-A6450), MODERATE/HIGH COMPRESSION BANDAGE (A6451, A6452), SELF-ADHERENT BANDAGE (A6453-A6455), CONFORMING BANDAGE (A6442-A6447), PADDING BANDAGE (A6441):

Most compression bandages are reusable. Usual frequency of replacement would be no more than one per week unless they are part of a multi-layer compression bandage system.

Conforming bandage dressing change is determined by the frequency of change of the selected underlying dressing.

GRADIENT COMPRESSION WRAP (A6545):

Coverage of a non-elastic gradient compression wrap (A6545) is limited to one per 6 months per leg. Quantities exceeding this amount will be denied as not reasonable and necessary. Refer to Policy Article for statement concerning non-coverage if the ulcer has healed.

▼ HCPCS Level II Codes and Description

A4450 TAPE, NON-WATERPROOF, PER 18 SQUARE INCHES

A4452 TAPE, WATERPROOF, PER 18 SQUARE INCHES

A4461 SURGICAL DRESSING HOLDER, NON-REUSABLE, EACH

A4463 SURGICAL DRESSING HOLDER, REUSABLE, EACH

A4465 NON-ELASTIC BINDER FOR EXTREMITY

A4649 SURGICAL SUPPLY; MISCELLANEOUS

A6010 COLLAGEN BASED WOUND FILLER, DRY FORM, STERILE, PER GRAM OF COLLAGEN

A6011 COLLAGEN BASED WOUND FILLER, GEL/PASTE, STERILE, PER GRAM OF COLLAGEN

A6021 COLLAGEN DRESSING, STERILE, PAD SIZE 16 SQ. IN. OR LESS, EACH

A6022 COLLAGEN DRESSING, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., EACH

A6023 COLLAGEN DRESSING, STERILE, PAD SIZE MORE THAN 48 SQ. IN., EACH

- A6024 COLLAGEN DRESSING WOUND FILLER, STERILE, PER 6 INCHES
- A6025 GEL SHEET FOR DERMAL OR EPIDERMAL APPLICATION, (E.G., SILICONE, HYDROGEL, OTHER), EACH
- A6154 WOUND POUCH, EACH
- A6196 ALGINATE OR OTHER FIBER GELLING DRESSING, WOUND COVER, STERILE, PAD SIZE 16 SQ. IN. OR LESS, EACH DRESSING
- A6197 ALGINATE OR OTHER FIBER GELLING DRESSING, STERILE, WOUND COVER, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., EACH DRESSING
- A6198 ALGINATE OR OTHER FIBER GELLING DRESSING, STERILE, WOUND COVER, PAD SIZE MORE THAN 48 SQ. IN., EACH DRESSING
- A6199 ALGINATE OR OTHER FIBER GELLING DRESSING, STERILE, WOUND FILLER, PER 6 INCHES
- A6203 COMPOSITE DRESSING, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6204 COMPOSITE DRESSING, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6205 COMPOSITE DRESSING, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6206 CONTACT LAYER, STERILE, 16 SQ. IN. OR LESS, EACH DRESSING
- A6207 CONTACT LAYER, STERILE, MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., EACH DRESSING
- A6208 CONTACT LAYER, STERILE, MORE THAN 48 SQ. IN., EACH DRESSING
- A6209 FOAM DRESSING, WOUND COVER, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6210 FOAM DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6211 FOAM DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6212 FOAM DRESSING, WOUND COVER, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6213 FOAM DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITH

ANY SIZE ADHESIVE BORDER, EACH DRESSING

A6214 FOAM DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING

A6215 FOAM DRESSING, WOUND FILLER, STERILE, PER GRAM

A6216 GAUZE, NON-IMPREGNATED, NON-STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITHOUT ADHESIVE BORDER, EACH DRESSING

A6217 GAUZE, NON-IMPREGNATED, NON-STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING

A6218 GAUZE, NON-IMPREGNATED, NON-STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING

A6219 GAUZE, NON-IMPREGNATED, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING

A6220 GAUZE, NON-IMPREGNATED, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING

A6221 GAUZE, NON-IMPREGNATED, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING

A6222 GAUZE, IMPREGNATED WITH OTHER THAN WATER, NORMAL SALINE, OR HYDROGEL, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITHOUT ADHESIVE BORDER, EACH DRESSING

A6223 GAUZE, IMPREGNATED WITH OTHER THAN WATER, NORMAL SALINE, OR HYDROGEL, STERILE, PAD SIZE MORE THAN 16 SQUARE INCHES, BUT LESS THAN OR EQUAL TO 48 SQUARE INCHES, WITHOUT ADHESIVE BORDER, EACH DRESSING

A6224 GAUZE, IMPREGNATED WITH OTHER THAN WATER, NORMAL SALINE, OR HYDROGEL, STERILE, PAD SIZE MORE THAN 48 SQUARE INCHES, WITHOUT ADHESIVE BORDER, EACH DRESSING

A6228 GAUZE, IMPREGNATED, WATER OR NORMAL SALINE, PAD SIZE 16 SQ. IN. OR LESS, STERILE, WITHOUT ADHESIVE BORDER, EACH DRESSING

A6229 GAUZE, IMPREGNATED, WATER OR NORMAL SALINE, STERILE, PAD SIZE MORE THAT 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING

A6230 GAUZE, IMPREGNATED, WATER OR NORMAL SALINE, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING

- A6231 GAUZE, IMPREGNATED, HYDROGEL, FOR DIRECT WOUND CONTACT, STERILE, PAD SIZE 16 SQ. IN. OR LESS, EACH DRESSING
- A6232 GAUZE, IMPREGNATED, HYDROGEL, FOR DIRECT WOUND CONTACT, STERILE, PAD SIZE GREATER THAN 16 SQ. IN., BUT LESS THAN OR EQUAL TO 48 SQ. IN., EACH DRESSING
- A6233 GAUZE, IMPREGNATED, HYDROGEL FOR DIRECT WOUND CONTACT, STERILE, PAD SIZE MORE THAN 48 SQ. IN., EACH DRESSING
- A6234 HYDROCOLLOID DRESSING, WOUND COVER, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6235 HYDROCOLLOID DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6236 HYDROCOLLOID DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6237 HYDROCOLLOID DRESSING, WOUND COVER, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6238 HYDROCOLLOID DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6239 HYDROCOLLOID DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6240 HYDROCOLLOID DRESSING, WOUND FILLER, PASTE, STERILE, PER FLUID OUNCE
- A6241 HYDROCOLLOID DRESSING, WOUND FILLER, DRY FORM, STERILE, PER GRAM
- A6242 HYDROGEL DRESSING, WOUND COVER, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6243 HYDROGEL DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6244 HYDROGEL DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING

- A6245 HYDROGEL DRESSING, WOUND COVER, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6246 HYDROGEL DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6247 HYDROGEL DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6248 HYDROGEL DRESSING, WOUND FILLER, GEL, STERILE, PER FLUID OUNCE
- A6250 SKIN SEALANTS, PROTECTANTS, MOISTURIZERS, OINTMENTS, ANY TYPE, ANY SIZE
- A6251 SPECIALTY ABSORPTIVE DRESSING, WOUND COVER, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6252 SPECIALTY ABSORPTIVE DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6253 SPECIALTY ABSORPTIVE DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING
- A6254 SPECIALTY ABSORPTIVE DRESSING, WOUND COVER, PAD SIZE 16 SQ. IN. OR LESS, WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6255 SPECIALTY ABSORPTIVE DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6256 SPECIALTY ABSORPTIVE DRESSING, WOUND COVER, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITH ANY SIZE ADHESIVE BORDER, EACH DRESSING
- A6257 TRANSPARENT FILM, STERILE, 16 SQ. IN. OR LESS, EACH DRESSING
- A6258 TRANSPARENT FILM, STERILE, MORE THAN 16 SQ. IN. BUT LESS THAN OR EQUAL TO 48 SQ. IN., EACH DRESSING
- A6259 TRANSPARENT FILM, STERILE, MORE THAN 48 SQ. IN., EACH DRESSING
- A6260 WOUND CLEANSERS, STERILE, ANY TYPE, ANY SIZE

A6261 WOUND FILLER, GEL/PASTE, STERILE, PER FLUID OUNCE, NOT OTHERWISE SPECIFIED.

A6262 WOUND FILLER, DRY FORM, STERILE, PER GRAM, NOT OTHERWISE SPECIFIED.

A6266 GAUZE, IMPREGNATED, OTHER THAN WATER, NORMAL SALINE, OR ZINC PASTE, STERILE, ANY WIDTH, PER LINEAR YARD

A6402 GAUZE, NON-IMPREGNATED, STERILE, PAD SIZE 16 SQ. IN. OR LESS, WITHOUT ADHESIVE BORDER, EACH DRESSING

A6403 GAUZE, NON-IMPREGNATED, STERILE, PAD SIZE MORE THAN 16 SQ. IN. LESS THAN OR EQUAL TO 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING

A6404 GAUZE, NON-IMPREGNATED, STERILE, PAD SIZE MORE THAN 48 SQ. IN., WITHOUT ADHESIVE BORDER, EACH DRESSING

A6407 PACKING STRIPS, NON-IMPREGNATED, STERILE, UP TO 2 INCHES IN WIDTH, PER LINEAR YARD

A6410 EYE PAD, STERILE, EACH

A6411 EYE PAD, NON-STERILE, EACH

A6412 EYE PATCH, OCCLUSIVE, EACH

A6413 ADHESIVE BANDAGE, FIRST AID TYPE, ANY SIZE, EACH

A6441 PADDING BANDAGE, NON-ELASTIC, NON-WOVEN/NON-KNITTED, WIDTH GREATER THAN OR EQUAL TO THREE INCHES AND LESS THAN FIVE INCHES, PER YARD

A6442 CONFORMING BANDAGE, NON-ELASTIC, KNITTED/WOVEN, NON-STERILE, WIDTH LESS THAN THREE INCHES, PER YARD

A6443 CONFORMING BANDAGE, NON-ELASTIC, KNITTED/WOVEN, NON-STERILE, WIDTH GREATER THAN OR EQUAL TO THREE INCHES AND LESS THAN FIVE INCHES, PER YARD

A6444 CONFORMING BANDAGE, NON-ELASTIC, KNITTED/WOVEN, NON-STERILE, WIDTH GREATER THAN OR EQUAL TO 5 INCHES, PER YARD

A6445 CONFORMING BANDAGE, NON-ELASTIC, KNITTED/WOVEN, STERILE, WIDTH LESS THAN THREE INCHES, PER YARD

A6446 CONFORMING BANDAGE, NON-ELASTIC, KNITTED/WOVEN, STERILE, WIDTH GREATER THAN OR EQUAL TO THREE INCHES AND LESS THAN FIVE INCHES, PER YARD

A6447 CONFORMING BANDAGE, NON-ELASTIC, KNITTED/WOVEN, STERILE, WIDTH GREATER THAN OR EQUAL TO FIVE INCHES, PER

YARD

A6448 LIGHT COMPRESSION BANDAGE, ELASTIC, KNITTED/WOVEN, WIDTH LESS THAN THREE INCHES, PER YARD

A6449 LIGHT COMPRESSION BANDAGE, ELASTIC, KNITTED/WOVEN, WIDTH GREATER THAN OR EQUAL TO THREE INCHES AND LESS THAN FIVE INCHES, PER YARD

A6450 LIGHT COMPRESSION BANDAGE, ELASTIC, KNITTED/WOVEN, WIDTH GREATER THAN OR EQUAL TO FIVE INCHES, PER YARD

A6451 MODERATE COMPRESSION BANDAGE, ELASTIC, KNITTED/WOVEN, LOAD RESISTANCE OF 1.25 TO 1.34 FOOT POUNDS AT 50% MAXIMUM STRETCH, WIDTH GREATER THAN OR EQUAL TO THREE INCHES AND LESS THAN FIVE INCHES, PER YARD

A6452 HIGH COMPRESSION BANDAGE, ELASTIC, KNITTED/WOVEN, LOAD RESISTANCE GREATER THAN OR EQUAL TO 1.35 FOOT POUNDS AT 50% MAXIMUM STRETCH, WIDTH GREATER THAN OR EQUAL TO THREE INCHES AND LESS THAN FIVE INCHES, PER YARD

A6453 SELF-ADHERENT BANDAGE, ELASTIC, NON-KNITTED/NON-WOVEN, WIDTH LESS THAN THREE INCHES, PER YARD

A6454 SELF-ADHERENT BANDAGE, ELASTIC, NON-KNITTED/NON-WOVEN, WIDTH GREATER THAN OR EQUAL TO THREE INCHES AND LESS THAN FIVE INCHES, PER YARD

A6455 SELF-ADHERENT BANDAGE, ELASTIC, NON-KNITTED/NON-WOVEN, WIDTH GREATER THAN OR EQUAL TO FIVE INCHES, PER YARD

A6456 ZINC PASTE IMPREGNATED BANDAGE, NON-ELASTIC, KNITTED/WOVEN, WIDTH GREATER THAN OR EQUAL TO THREE INCHES AND LESS THAN FIVE INCHES, PER YARD

A6457 TUBULAR DRESSING WITH OR WITHOUT ELASTIC, ANY WIDTH, PER LINEAR YARD

Documentation Requirement

1. Compression wrap is limited to one per 6 months per leg. Quantities exceeding this amount will be denied as not reasonable and necessary.
2. The treating physician's order must specify (a) the type of dressing (e.g., hydrocolloid wound cover, hydrogel wound filler, etc.), (b) the size of the dressing (if appropriate), (c) the number/amount to be used at one time (if more than one), (d) the frequency of dressing change, and (e) the expected duration of need.
3. A new order is needed if a new dressing is added or if the quantity of an existing dressing to be used is increased. A new order is not routinely needed if the quantity of dressings used is decreased. However, a new order is required at least every 3 months for each dressing being used even if the quantity used has remained the same or decreased.
4. Information defining the number of surgical/debrided wounds being treated with a dressing, the reason for dressing use (e.g., surgical wound, debrided wound, etc.), and whether the dressing is being used as a primary or secondary dressing or for some non covered use (e.g., using a dressing for wound cleansing) must be obtained from the physician or home care nurse. The source of that information and date obtained must be documented and sent in with the claim.
5. Current clinical information which supports the reasonableness and necessity of the type and quantity of surgical dressings provided must be documented and sent in with the claim. Evaluation of a Member's wound(s) must be performed at least on a monthly basis unless there is documentation sent in with the claim which justifies why an evaluation could not be done within this timeframe and what other monitoring methods were used to evaluate the Member's need for dressings. Evaluation is expected on a more frequent basis (e.g., weekly) in Members with heavily draining or infected wounds. The evaluation may be performed by a nurse, physician or other health care professional. This evaluation must include the type of each wound (e.g., surgical wound, pressure ulcer, burn, etc), its location, its size (length x width in cm.) and depth, the amount of drainage, and any other relevant information. This information must be documented and sent in with the claim.
6. When codes A4649, A6261 or A6262 are billed, the claim must include a narrative description of the item (including size of the product provided), the manufacturer, the brand name or number, and information justifying the medical necessity for the item. This information must be entered in the narrative field of the electronic claim.

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. This bottoming out criterion should be tested with the patient

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 sidelying position.

**The staging of pressure ulcers used in this policy is as follows
(National Pressure Ulcer Advisory Panel, 2016 Revision):**

Stage 1 Pressure Injury: Non-blanchable erythema of intact skin
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 Pressure Injury: Partial-thickness skin loss with exposed dermis
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 (MASD) including incontinence associated dermatitis (IAD), intertriginous dermatitis (ITD), medical adhesive related skin injury (MARS), or traumatic wounds (skin tears, burns, abrasions).

Stage 3 Pressure Injury: Full-thickness skin loss
Full-thickness loss of skin, in which adipose (fat) is visible in the ulcer and granulation tissue and epibole (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 Pressure Injury: Full-thickness skin and tissue loss
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. Epibole (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.

Unstageable Pressure Injury: Obscured full-thickness skin and tissue loss
Full-thickness skin and tissue loss in which the extent of tissue damage within the ulcer cannot be confirmed because it is obscured by slough or eschar. If slough or eschar is removed, a Stage 3 or Stage 4 pressure

injury will be revealed. Stable eschar (i.e. dry, adherent, intact without erythema or fluctuance) on the heel or ischemic limb should not be softened or removed.

Deep Tissue Pressure Injury: Persistent non-blanchable deep red, maroon or purple discoloration Intact or non-intact skin with localized area of persistent non-blanchable deep red, maroon, purple discoloration or epidermal separation revealing a dark wound bed or blood filled blister. Pain and temperature change often precede skin color changes. Discoloration may appear differently in darkly pigmented skin. This injury results from intense and/or prolonged pressure and shear forces at the bone-muscle interface. The wound may evolve rapidly to reveal the actual extent of tissue injury, or may resolve without tissue loss. If necrotic tissue, subcutaneous tissue, granulation tissue, fascia, muscle or other underlying structures are visible, this indicates a full thickness pressure injury (Unstageable, Stage 3 or Stage 4). Do not use DTPI to describe vascular, traumatic, neuropathic, or dermatologic conditions.

▼ Definitions

Debridement of a wound may be any type of debridement (examples given are not all-inclusive):

1. Surgical (e.g. sharp instrument or laser)
2. Mechanical (e.g. irrigation or wet-to-dry dressings)
3. Chemical (e.g. topical application of enzymes)
4. Autolytic (e.g. application of occlusive dressings to an open wound)

▼ Important Note:

Northwood's Medical Policies are developed to assist Northwood in administering plan benefits and determining whether a particular DMEPOS product or service is reasonable and necessary. Equipment that is used primarily and customarily for a non-medical purpose is not considered durable medical equipment.

Coverage determinations are made on a case-by-case basis and are subject to all of the terms, conditions, limitations, and exclusions of the member's contract including medical necessity requirements.

The conclusion that a DMEPOS product or service is reasonable and necessary does not constitute coverage. The member's contract defines which DMEPOS product or service is covered, excluded or limited. The policies provide for clearly written, reasonable and current criteria that have been approved by Northwood's Medical Director.

The clinical criteria and medical policies provide guidelines for determining the medical necessity for specific DMEPOS products or services. In all cases, final benefit determinations are based on the applicable contract language. To the extent there are any conflicts between medical policy guidelines and applicable contract language, the contract language prevails. Medical policy is not intended to override the policy that defines the member's benefits, nor is it intended to dictate to providers how to direct care. Northwood Medical policies shall not be interpreted to limit the benefits afforded to Medicare or Medicaid members by law and regulation and Northwood will use the applicable state requirements to determine required quantity limit guidelines.

Northwood's policies do not constitute medical advice. Northwood does not provide or recommend treatment to members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

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Applicable URAC Standard

Core 8	Staff operational tools and support
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Change/Authorization History

Revision Number	Date	Description of Change	Prepared / Reviewed by	Approved by	Review Date:
A	11-20-06	Initial Release	Rosanne Brugnani	Ken Fasse	n/a
01	12-2008	Changed descriptions for the staging of pressure ulcers and added HCPC codes A6413 and	Susan Glomb	Ken Fasse	n/a

		A6545			
02		Annual Review – no additional changes	Susan Glomb	Ken Fasse	Dec.2008
03	01/01/09	Added A6545 Revised A6010-A6024 A6196-A6199 A6203-A6215 A6219-A6248 A6251-A6266 A6407 Codes were changed to reflect items as “sterile”	Susan Glomb	Ken Fasse	
04	12-22-09	Annual Review/ no changes	Susan Glomb	Ken Fasse	Dec.2009
05	01-05-10	Narrative change: A6549. Gradient compression stocking/sleeve, not otherwise specified. Discontinued codes: A6200; Composite dressing, pad size 16 sq. in. or less, without adhesive border, each dressing. Now A6251. A6201; Composite dressing, pad size more than 16 sq. in. but less than or equal to 48 sq.in., without adhesive border, each dressing. Now A6252. A6202; Composite dressing, pad size more than 48 sq.in. without adhesive border, each dressing. Now A6253. A6542; Gradient compression stocking, custom made. Now A6549. A6543; Gradient compression stocking, lymphedema . Now A6549	Susan Glomb	Ken Fasse	
06	12-03-10	Annual Review – No Changes	Susan Glomb	Ken Fasse	Dec.2010
07	07-20-11	Added Important Note to all Medical Policies	Susan Glomb	Dr. B. Almasri	
08	11-7-11	Annual Review. Policy References Added. Inclusion of Abdominal Binder/Abdominal Support into this Policy.	Susan Glomb	Dr. B. Almasri	November 2011
09	12-04-12	Annual review – no changes.	Susan Glomb	Dr. B. Almasri	Dec. 2012
10	12-30-13	Annual review. No changes.	Susan Glomb	Dr. B. Almasri	
11	12-1-14	Annual Review. No changes	Susan Glomb	Dr. B. Almasri	

12	12-11-15	Annual review. Updated policy with Medicare policy. Updated references.	Susan Glomb	Dr. B. Almasri	
13	5-24-16	Clarified # 4- added that dressings are not covered for use in wound cleansing.	Susan Glomb	Dr. B. Almasri	
14	12-08-16	Annual Review. No Changes.	Lisa Wojno	Dr. B. Almasri	December 2016
15	12-16-17	Annual review. Revised: Staging guidelines to reflect NPUAP 2016 Staging Consensus Conference.	Carol Dimech	Dr. C. Lerchin	December 2017
16	12-7-18	Annual review. Medicare references updated.	Carol Dimech	Dr. C. Lerchin	December 2018