

McKESSON

Collagen Dressings

ADVANCED WOUND DRESSING | STERILE

Intended for management of full and partial thickness wounds.

PRODUCT AVAILABILITY

MFR #	SIZE	AVAILABILITY
16-1850	2 in x 2 in (5.1 cm x 5.1 cm)	10 DRESSINGS PER BOX
16-1852	4 in x 4 in (10.2 cm x 10.2 cm)	10 DRESSINGS PER BOX

DESCRIPTION

The McKesson Collagen Dressing is an advanced wound dressing made from collagen, sodium alginate, carboxymethylcellulose (CMC), and ethylenediaminetetraacetic acid (EDTA). The McKesson Collagen Dressing is a topically applied wound dressing that will transform into a soft gel sheet when in contact with wound exudates. The dressing maintains a moist wound environment that aids in the formation of granulation tissue and epithelialization. The EDTA in the dressing removes zinc to inhibit the activity of matrix metalloproteinases (MMP), thus creating a suitable environment for wound healing. McKesson Collagen Dressings may be trimmed and layered for the management of deep wounds.

McKESSON

Questions? Call 1-800-777-4908

Satisfaction Guaranteed

If you are not completely satisfied with any McKesson Brands product, you may return it for a full refund or credit.

Distributed by McKesson Medical-Surgical Inc.
Richmond, VA 23233
PVN A0117
Made in USA

INDICATIONS FOR USE

McKesson Collagen Dressings are indicated for management of full and partial thickness wounds including: pressure ulcers, diabetic ulcers, ulcers caused by mixed vascular aetiologies, venous ulcers, donor and graft sites, abrasions, traumatic wounds healing by secondary intention, dehisced surgical wounds, first and secondary degree burns.

CONTRAINDICATIONS

McKesson Collagen Dressings should not be used on patients with a known allergy or sensitivity to animal-derived collagen, any other ingredient or on third degree burns.

PRECAUTIONS

If sensitivity to McKesson Collagen Dressing develops, discontinue use. McKesson Collagen Dressings may be used under compression therapy under the supervision of a health care professional.

DIRECTIONS FOR USE

1. Debride and irrigate the wound bed with an appropriate wound cleanser or distilled water before application.
2. Cut the dressing to fit the exact wound size. For heavily exuding wounds, apply the dressing directly to the wound bed. For dry wounds with very little exudate, moisten the wound bed with distilled water to begin the gelling process.
3. Cover the McKesson Collagen Dressing with an appropriate secondary dressing.
4. Change dressings daily or as good nursing practice dictates.

CARE AND REMOVAL





1. Remove the secondary dressing with care. Gently remove McKesson Collagen Dressing ensuring that the dressing removal does not damage any newly formed tissues.
2. Use an appropriate wound cleanser or distilled water to cleanse the wound site prior to applying a new McKesson Collagen Matrix Dressing as per application instructions.

NOTE: As with any medical device, always refer to the full instructions for use prior to application.

CAUTION: Federal (USA) law restricts this device to sale by or on the order of a physician.

Store at 50–86°F (10–30°C). Avoid storage in direct sunlight/ fluorescent lighting and store in a cool, dry, and well-ventilated area. Not made with natural rubber latex. Contents STERILE in unopened, undamaged inner package.

SYMBOL LEGEND

SYMBOL	TITLE	STANDARD / REFERENCE #	DESCRIPTION
	Sterilized using irradiation	ISO 15223-1; 5.2.4	Indicates a medical device that has been sterilized using irradiation.
	Caution	ISO 15223-1; 5.4.4	Indicates the need for the user to consult the instructions for use for important cautionary information such as warnings and precautions that cannot, for a variety of reasons, be presented on the medical device itself.
	The symbol statement for Prescription Device	21 CFR 801.109; N/A	Indicates that the product is a medical device as defined in 21 CFR 820.3(l) and Federal Law (USA) restricts this device to sale by or on the order of a physician [21 CFR 801.109]
 DO NOT REUSE	Do not re-use	ISO 15223-1; 5.4.2	Indicates a medical device that is intended for one use, or for use on a single patient during a single procedure.