MEDIHONEY®
Barrier Cream

Barrier Cream

MediHoney® Barrier Cream is produced with medical-grade antibacterial Manuka honey.

30%
ANTIBACTERIAL MANUKA HONEY*

* by weight

INTEGRA®
LIMIT UNCERTAINTY

MEDIHONEY®
Barrier Cream

Product Code	Product Description	Size	Qty/Box	PIP Code	NHS Code
582	Barrier Cream	50g	1	338-7644	ELY289
800	Barrier Cream Sachets	2g	20	369-1276	ELY374

Features and Benefits

MediHoney Barrier Cream can be applied to intact and at-risk skin to provide protection from bodily fluids and moisture-associated skin breakdown.

- Made with medical-grade antibacterial Manuka honey. Contains no parabens, mineral oils, lanolin, steroids, added fragrance or colour.
- Medical-grade Manuka honey delivers broad spectrum antimicrobial activity.
- Medical-grade Manuka honey has a low pH of 3.5-4.5. Lowering pH has been associated with wound healing.
- Formulated to provide a dry to touch layer to protect skin from damage caused by friction and shear, allowing adhesive dressings to be used effectively.
- Designed to dry quickly, allowing the skin to remain breathable, which helps to prevent maceration.
- Has been safely used on patients of all ages.
- Protects skin at risk for moisture damage.
- Helps prevent damage to skin caused by frequent washing: MediHoney Barrier Cream does not need to be removed prior to reapplication.
- Available in 2g sachet and 50g tube to meet hospital and community needs.

Indications

- Protects at-risk skin from breakdown associated with incontinence.
- Protects at-risk skin folds from moisture associated skin irritation and breakdown.
- Can be used around wound edges (peri-wound) to protect at-risk skin from irritation or breakdown associated with moisture from wound exudate.
- Helps prevent maceration.

Contraindication

- MediHoney Barrier Cream is contraindicated for patients that have a known sensitivity to honey.

Availability of these products might vary from a given country or region to another, as a result of specific local regulatory approval or clearance requirements for sale in such country or region.
- Non contractual document. The manufacturer reserves the right, without prior notice, to modify the products in order to improve their quality.
- Consult product labels and inserts for any indication, contraindications, hazards, warnings, precautions, and instructions for use.

Additional information for EMEA Customers only:
Products mentioned in this document are CE class I, IIa, IIb or III devices. Contact Integra should you need any additional information on devices classification. All the medical devices mentioned on this document are CE marked according to European council directive 93/42/EEC on medical devices and its relatives, unless specifically identified as "NOT CE MARKED".

For more information or to place an order, please contact:

Customer Service
United Kingdom +44 (0) 1264 312 725 ▪ +44 (0) 1264 312 821 fax
integralife.eu

CE

Manufacturer:

Derma Sciences, Inc.
104 Shorting Road
Toronto, Ontario M1S 3S4 ▪ Canada

EC REP Scanlan Group B.V.
Postbus 75664
Schiphol Triport 1118 ZS
The Netherlands
Int: +31 20 653-0553