

MANUKA HONEY EXTRA STRENGTH- allantoin gel
First Honey LLC

Manuka Honey Ointment Extra Strength

Drug Facts

Active ingredient

Allantoin 0.5%

Purpose

Skin Protectant

Uses

- Temporarily protects minor
- cuts
- scrapes
- burns
- helps prevent and temporarily protects and helps relieve chafed, chapped, or cracked skin

Warnings

For external use only

Do not use

- In the eyes
- On deep or puncture wounds
- Animal bites
- Serious burns

Stop use and ask a doctor if

- Conditions worsen
- Symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

If swallowed, get medical help, or contact a Poison Control Center right away.

Directions

Apply as needed.

Inactive ingredients

Helianthus Annuus Extract, Lecithin, Mānuka Honey, Oryza Sativa Bran Extract, Propanediol, Potassium Sorbate, Rosmarinus Officinalis Leaf Extract, Sodium Acrylates Copolymer, Sodium Benzoate, Tetrasodium Glutamate Diacetate, Tocopherol, Water (Aqua).

Questions?

1-615-488-8168

Package Labeling:

Healing should be sweet.

firsthoney

firsthoney

Mānuka Honey Ointment
EXTRA STRENGTH

Net Weight
0.75oz – 21g

First Honey is the global leader in over-the-counter medical honey treatments for natural, antibiotic free wound care. Our team of beekeepers and scientists have dedicated themselves to providing the highest quality honey, harvested from the powerful nectar of New Zealand's native Mānuka flower.

Drug Facts

Active ingredient Purpose
Allantoin 0.5% Skin Protectant

Uses ■ Temporarily protects minor ■ cuts ■ scrapes ■ burns ■ helps prevent and temporarily protects and helps relieve chafed, chapped, or cracked skin

Warnings For external use only

Do not use ■ In the eyes ■ On deep or puncture wounds ■ Animal bites ■ Serious burns

Stop use and ask a doctor if ■ Conditions worsen ■ Symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of children. If swallowed, get medical help, or contact a Poison Control Center right away.

Directions Apply as needed.

Inactive ingredients
Helianthus Annuus Extract, Lecithin, Mānuka Honey, Oryza Sativa Bran Extract, Propanediol, Potassium Sorbate, Rosmarinus Officinalis Leaf Extract, Sodium Acrylates Copolymer, Sodium Benzoate, Tetrasodium

Directions for use:
Clean the affected area. Apply ointment directly to wound. Cover with a sterile bandage.

Dressing change:
Depending on wound type, the product may require a daily application. Re-apply as necessary.

Contraindications:
Do not use on patients with known sensitivity to honey. Minor stinging may be experienced due to osmotic action and/or the low pH of honey. This should be temporary, if pain persists, discontinue use of the dressing and gently irrigate the wound with sterile saline solution.

Caution: Although there is no record of increased blood sugar levels in patients with diabetes due to the use of honey

Glutamate Diacetate,
Tocopherol, Water (Aqua).

Questions?
1-615-488-8168

Distributed by First Honey
LLC, 18 Fairfield Avenue
Nashville, TN 37210
firsthoney.com

NDC 81995-050-01

dressings, it is advisable
to monitor the levels during
use. Discontinue use after
30 days, and seek medical
help if the wound has
not healed.

Made in New Zealand.

**MEDICAL
GRADE
MĀNUKA
HONEY**

Created for the moments
in everyday life when you
need safe and effective
care for the ones you love.

firsthoney.com
@firsthoneynz



LOT



AW0050C01

8 54056 00835 6

MANUKA HONEY EXTRA STRENGTH

allantoin gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81995-050
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PROPANEDIOL (UNII: 5965N8W85T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
ROSEMARY (UNII: IJ67X351P9)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
TOCOPHEROL (UNII: R0ZB2556P8)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81995-050-01	1 in 1 CARTON	09/01/2024	
1		21 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	09/01/2024	

Labeler - First Honey LLC (080994597)

Revised: 2/2025

First Honey LLC