ALLANTOIN- advanced derma therapy spray
Harmon Store Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Derma Therapy Spray

Active ingredient

Allantoin 0.5%

Purpose

Skin Protectant

Uses

Temporarily protects and helps relieve chapped or cracked skin

Warnings

For external use only.

Contents under pressure. Do not puncture or incinerate. Do not store above 120 °F. Intentional misuse by deliberately concentrating and inhaling contents can be harmful or fatal.

Do not use

- on deep or puncture wounds
- on animal bites
- on serious burns

When using this product

- avoid spraying in eyes

Stop use and ask a doctor if

- condition worsens
- 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

- shake well before using
- apply as needed
- spray product onto affected area(s) and massage into skin in a circular motion
- for scars on face or near eyes spray product into palm of hand and apply with fingertips

Other information

store at room temperature

Inactive ingredients

Water, Dimethyl Ether, Isostearyl Isostearate, Isopropyl Isostearate, PPG-3 Benzyl Ether Myristate, Pentaerythrityl Tetracaprylate/Caprate, Methylsilanol Hydroxyproline Aspartate, Glycerin, Butylene Glycol, Allium Cepa (Onion) Bulb Extract, Sucrose Palmitate, Glyceryl Stearate, Glyceryl Stearate Citrate, Sucrose, Mannan, Xanthan Gum, Cetyl Hydroxyethylcellulose, Rutin, Palmitoyl Tripeptide-1, Palmitoyl Tetrapeptide-7, Phaseolus Lunatus (Green Bean) SEed Extract, Phenoxyethanol, Fragrance, Potassium Sorbate, Salicylic Acid

Questions?

Call 1-866-964-0939

Principal Display Panel

CoreValue

Advanced

Derma Therapy

Spray

Allantoin 0.5% - Skin Protectant

- Improves overall appearance of scars*
- Improves the appearance of texture and smoothness on stretch marks*

Dermatologist Tested

Paraben-Free

Shake well before using.

Net WT 3 OZ (85 g)

ALLANTOIN

advanced derma therapy spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63940-022
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.425 g in 85 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
DIMETHYL ETHER (UNII: AM13FS69BX)	
ISOSTEARYL ISOSTEARATE (UNII: IV0Z586Z4Y)	
PPG-3 BENZYL ETHER MYRISTATE (UNII: 8075L58MKO)	
PENTAERYTHRITYL TETRACAPRYLATE/TETRACAPRATE (UNII: 832C4KF14X)	
GLYCERIN (UNII: PDC6A3C0OX)	

BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ONION (UNII: 492225Q21H)	
SUCROSE PALMITATE (UNII: 3OSQ643ZK5)	
GLYCERYL STEARATE CITRATE (UNII: WH8T92A065)	
SUCROSE (UNII: C151H8M554)	
YEAST MANNAN (UNII: 91R887N59P)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CETYL HYDROXYETHYLCELLULOSE (350000 MW) (UNII: T7SWE4S2TT)	
RUTIN (UNII: 5G06TVY3R7)	
PALMITOYL TRIPEPTIDE-1 (UNII: RV743D216M)	
PALMITOYL TETRAPEPTIDE-7 (UNII: Q41S464P1R)	
LIMA BEAN (UNII: 112YH1ZMX2)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63940-022-03	85 g in 1 CAN; Type 0: Not a Combination Product	09/04/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	09/04/2015	

Labeler - Harmon Store Inc. (804085293)

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