MEDERMA ADVANCED SCAR- allantoin gel
HRA PHARMA AMERICA, INC.

MEDERMA® Advanced Scar Gel *DRUG FACTS*

Active Ingredients

Allantoin 0.5%

Purpose

Skin Protectant

Uses

- Temporarily protects and helps relieve chapped or cracked skin

Warnings

- For external use only

When using this product

- Do not get into 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

Do not use on

- Deep puncture wounds
- Animal bites
- Serious burns

Keep out of reach of children.

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

Directions

- Apply as needed

Other Information

- Store at room temperature

Inactive Ingredients

- Water, PEG-4, Alcohol, Xanthan Gum, Allium Cepa (Onion) Bulb Extract, Panthenol, Fragrance, Lecithin, Methylparaben, Sorbic Acid, Sodium Hyaluronate

Questions or Comments?

- For more information call 1-833-426-6733 or visit www.mederma.com

PRINCIPAL DISPLAY PANEL

MEDERMA® ADVANCED SCAR GEL

Clinically shown to visibly reduce the appearance of scars

No 1 Doctor & Pharmacist recommended scar brand

Skin Protectant

1 x GEL TUBE

NET WT. 1.76 Oz (50g)

Unique Triple-Action Formula penetrates into the skin

MEDERMA ADVANCED SCAR

allantoin gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73302-201
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
Water (UNII: 059QF0KO0R)	
Polyethylene Glycol 200 (UNII: R95B8J264J)	
Alcohol (UNII: 3K9958V90M)	
Xanthan Gum (UNII: TTV12P4NEE)	
Onion (UNII: 492225Q21H)	
Sorbic Acid (UNII: X045WJ989B)	
Panthenol (UNII: WV9CM0O67Z)	
Egg Phospholipids (UNII: 1Z74184RGV)	
Methylparaben (UNII: A2I8C7HI9T)	
Hyaluronate Sodium (UNII: YSE9PPT4TH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73302-201-20	1 in 1 BOX	11/01/2013	
1		20 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:73302-201-50	1 in 1 BOX	11/01/2013	
2		50 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	11/01/2013	

Labeler - HRA PHARMA AMERICA, INC. (081160441)

Revised: 11/2023 HRA PHARMA AMERICA, INC.