RITE AID SCAR- allantoin 0.5% gel Rite Aid

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.

Rite Aid Scar Gel

Allantoin 0.5%

Skin Protectant.

Temporarily protects and helps relieve chapped or cracked skin.

For external use only.

When using this product keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if the condition worsens or symptoms persist for more than 7 days or clear up and occur again within a few days.

Do not use on deep or puncture wounds, animal bites, and serious burns.

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

Apply as needed.

Water, Alcohol Denat., PEG-4, Propylene Glycol, Xanthan Gum, Allium Cepa (Onion) Bulk Extract, Panthenol, Sodium Hyaluronate, Aloe Barbadensis Leaf Extract, Chamomilla Recutita (Matricaria) Flower Extract, Cucumis Sativus (Cucumber) Fruit Extract, Tocopherol, Glycerin, Juglans Nigra (Black Walnut) Leaf Extract, Ethylhexylglycerin, Benzyl Alcohol, Fragrance.



RITE AID SCAR

allantoin 0.5% gel

Product Information

GLYCERIN (UNII: PDC6A3C0OX)

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-1297
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Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	0.5 a in 100 a

inactive ingredients		
Ingredient Name	Strength	
POLYETHYLENE GLYCOL 200 (UNII: R95B8J264J)		
TOCOPHEROL (UNII: R0ZB2556P8)		
PANTHENOL (UNII: WW9CM0O67Z)		
WATER (UNII: 059QF0KO0R)		

ONION (UNII: 492225Q21H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BLACK WALNUT (UNII: 02WM57RXZJ)	
CHAMOMILE (UNII: FGL3685T2X)	
CUCUMBER (UNII: YY7C30VXJT)	
ALCOHOL (UNII: 3K9958V90M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822- 1297-1	1 in 1 BOX	09/15/2022	
1		19.8 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 final	M016	09/15/2022	

Labeler - Rite Aid (014578892)

Registrant - Derma Care Research Labs, LLC (116817470)

Establishment				
Name	Address	ID/FEI	Business Operations	
Derma Care Research Labs, LLC		116817470	manufacture(11822-1297)	

Revised: 6/2023 Rite Aid