QC SCAR- allantoin 0.5% gel CDMA

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.

QC 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.



QC SCAR

allantoin 0.5% gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63868-271

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z) ALLANTOIN 0.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 200 (UNII: R95B8J264J)	
TOCOPHEROL (UNII: R0ZB2556P8)	
PANTHENOL (UNII: WV9CM0O67Z)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
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)	

F	Packaging						
#	tem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:63868-271- 01	1 in 1 BOX	08/16/2021				
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	part347	08/16/2021			

Labeler - CDMA (011920774)

Registrant - Derma Care Research Labs, LLC (116817470)

Establishment				
Name	Address	ID/FEI	Business Operations	
Derma Care Research Labs		116817470	manufacture(63868-271)	

Revised: 3/2023 CDMA