DCH SCAR- allantoin 0.5% gel
Derma Care Research Labs, LLC

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.

DCH Scar Gel, Allantoin 0.5%

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.

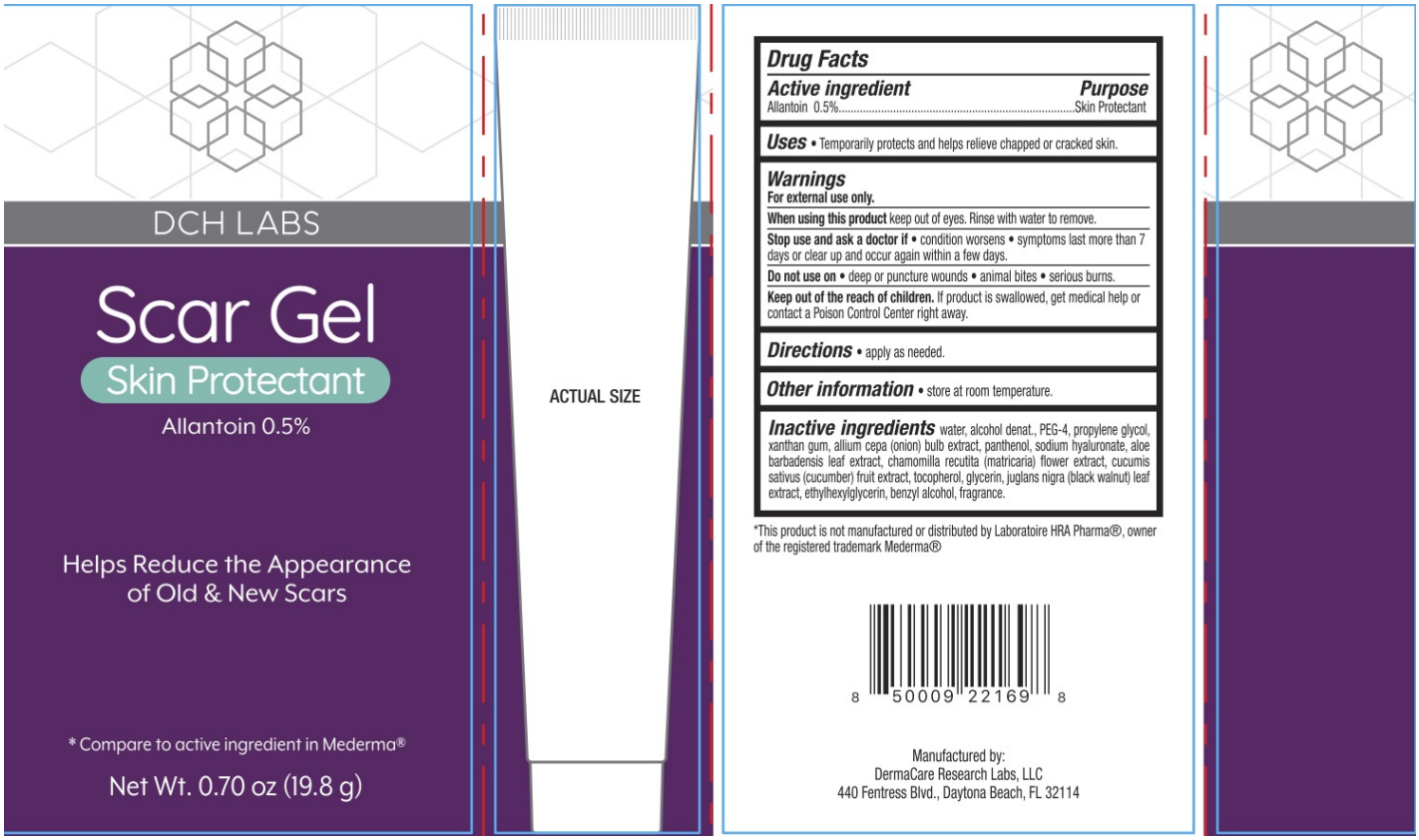
Stop use and ask a doctor if the condition worsens or symptoms persist for more than 7 days or clear up and occur again within a few days.

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.



DCH SCAR

allantoin 0.5% gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72839-097
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72839-097-01	1 in 1 BOX	10/05/2021	
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	10/05/2021	

Labeler - Derma Care Research Labs, LLC (116817470)

Registrant - Derma Care Research Labs, LLC (116817470)

Establishment

Name	Address	ID/FEI	Business Operations
Derma Care Research Labs		116817470	manufacture(72839-097)

Revised: 7/2023 Derma Care Research Labs, LLC