CVS HEALTH ADVANCED SCAR GEL- advanced scar gel gel CVS Health

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.

Scar Gel

Drug Facts

Active ingredients

Allantion 0.5%

Purpose

Skin Protectant

Uses

Temporarily protects and helps 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 for more than 7 days or clears up and occur again within a few days

Do Not Use on

- deep or puncture wounds
- animal bites
- serious burns

Keep out of the reach of children. If swallowed get medical help or contact a Poison Control Center immediately.

Directions

Apply as needed

Other information

Store at room temperature

Inactive ingredient

Water (Purified), PEG 300, Allium Cepa (Onion) Bulb Extract, Xanthan Gum, Alcohol, Methylparaben, Sorbic Acid, Panthenol, Sodium Hyaluronate, Fragrance

Principal Display Panel – Tube

Principal Display Panel – Tube

CVS Health NDC 69842-725-47

Advanced Scar Gel

0.5% Allantoin

NET WT. 1.76 oz (50g)



Advanced

car Ge

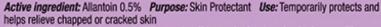
SPECIALLY FORMULATED FOR SCARS

SKIN PROTECTANT

Dermatologist tested

Specially formulated for scars caused by:
• Acne • Stretch marks • Injury • Burns • Surgery

NET WT 1.76 OZ (50 g)



Warnings: For external use only. When using this product • do not get into eyes. Stop use and ask a doctor if . condition worsens . symptoms persist for 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 immediately.

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Principal Display Panel - Carton

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CVS Health NDC 69842-725-47

Advance Scar Gel

Skin Protectant

NET WT. 1.76 oz (50g)





CVS HEALTH ADVANCED SCAR GEL

advanced scar gel gel

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:69842-725

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient NameBasis of StrengthStrengthALLANTO IN (UNII: 344S277G0 Z) (ALLANTO IN - UNII: 344S277G0 Z)ALLANTO IN5 mg in 1 g

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
POLOXAMER 407 (UNII: TUF2IVW3M2)				
ONION (UNII: 492225Q21H)				
XANTHAN GUM (UNII: TTV12P4NEE)				
ALCOHOL (UNII: 3K9958V90M)				
METHYLPARABEN (UNII: A218 C7H19 T)				
Sorbic Acid (UNII: X045WJ989B)				
PANTHENOL (UNII: WV9CM0O67Z)				
HYALURO NATE SO DIUM (UNII: YSE9 PPT4TH)				

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69842-725-47	1 in 1 CARTON	05/22/2019			
1		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 final	part347	05/22/2019				

Labeler - CVS Health (062312574)

Registrant - Sheffield Pharmaceuticals LLC (151177797)

Establishment						
Name	Address	ID/FEI	Business Operations			
Sheffield Pharmaceuticals LLC		151177797	MANUFACTURE(69842-725)			

Revised: 11/2019 CVS Health