OLP SCAR CREAM- allantoin cream
Ohio Lab Pharma

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.

Active Ingredients: Allantoin 0.5%

Keep out of reach of Children

Keep out of reach of Children: Keep out of reach of babies and children

Dosage and Administration: Gently massage into scar twice a day for 2-3 minutes

Purpose: Skin Protectant

Indication and Usage: Temporarily protects and helps relieve chapped or cracked skin.

Inactive Ingredients

- Cetostearyl alcohol- mineral oil- ceteareth 20- propylene glycol- methyl paraben- propylparaben- vitamin E- Stearic acid - trolamine
- RosmarinusOfficinalis (Rosemary)Extract/ Portulaca Oleracea Extract/Thymus Vulgaris Thyme) Leaf Extract/Jasminum Officinale (jasmine) Flower/ Leaf Extract?Chamomila) Recutita (Matricaria) Flower Water, Aureobasidium Pullulans Ferment Extract/ 1.2-Hexanedio

WARNING

Warning: 1. In case of having following symptoms after using this, you're advised to stop using it immediately. If you keep using it, the symptoms will get worse and need to consult a dermatologist. 1) In case of having problems such as red rash, swollenness, itching, stimulation during usage. 2) In case of having the same symptoms above on the part you put this product on by direct sunlight. 2. You are banned to use it on the part where you have a scar, eczema, or dermatitis. 3. In case of getting it into your eyes, you have to wash it immediately

Warnings

- In case of having the following symptoms after using this cream, you are advised to stop using it immediately. If you keep using it , the symptoms will get worse and need to consult a drematologist .1- in case of having problems such as red rash, swollllenness, itching, stimulation during usage. 2- in case of having the same symptoms above on the part you put this product on by direct sunlight
- you are banned to use it on the part where you have a eczema, lor dermatitis.
- in case of getting it into your eyes, you have to wash it immediately.

Questions

visit www.ohiolabpharma.us

Keep out of reach of children and babies

package label

Maximum strength

Net weight 0.7 oz (20g)

NDC#70648-333-01

OLP SCAR CREAM

allantoin cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70648-333
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**
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
EDETATE DISODIUM DIHYDRATE (UNII: 7FLD91C86K)	
MINERAL OIL (UNII: T5L8T28FGP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
THYMUS VULGARIS LEAF (UNII: GRX3499643)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
PORTULACA OLERACEA WHOLE (UNII: D5J3623SV2)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
ROSMARINUS OFFICINALIS WHOLE (UNII: EA3289138M)	
JASMINUM OFFICINALE FLOWER (UNII: 0Q8K841432)	
MATRICARIA CHAMOMILLA LEAF (UNII: 6I9LN466F0)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70648-333-01	1 in 1 CARTON	10/12/2017	
1		20 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	10/12/2017	

Labeler - Ohio Lab Pharma (080215854)

Establishment

Name	Address	ID/FEI	Business Operations
Ohio Lab Pharma		080215854	manufacture(70648-333)

Revised: 10/2017 Ohio Lab Pharma