DERMOVIX- allantoin,petrolatum patch
Binger Consulting Corporation

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.

Dermovix

Drug Facts

Active ingredient

Allantoin 2.00%

Petrolatum 30.00%

Purpose

Skin Protectant

Skin Protectant

Keep out of reach of children. If pregnant or breast feeding, contact physician prior to use.

Uses

• Scar Management • Temporarily protects minor cuts, scrapes and burns

Warnings

For external use only

Do not use

• on deep or puncture wounds, animal bites, serious burns

• in large quantities, particularly over raw surfaces or blistered areas

When using this product

• avoid contact with eyes

Stop use and ask a doctor if

• condition worsens symptoms last more than 7 days or clear up and occur again within a few days

Directions adults and children 12 years and over apply to affected area; change patch 1 to 2 times daily

How to apply

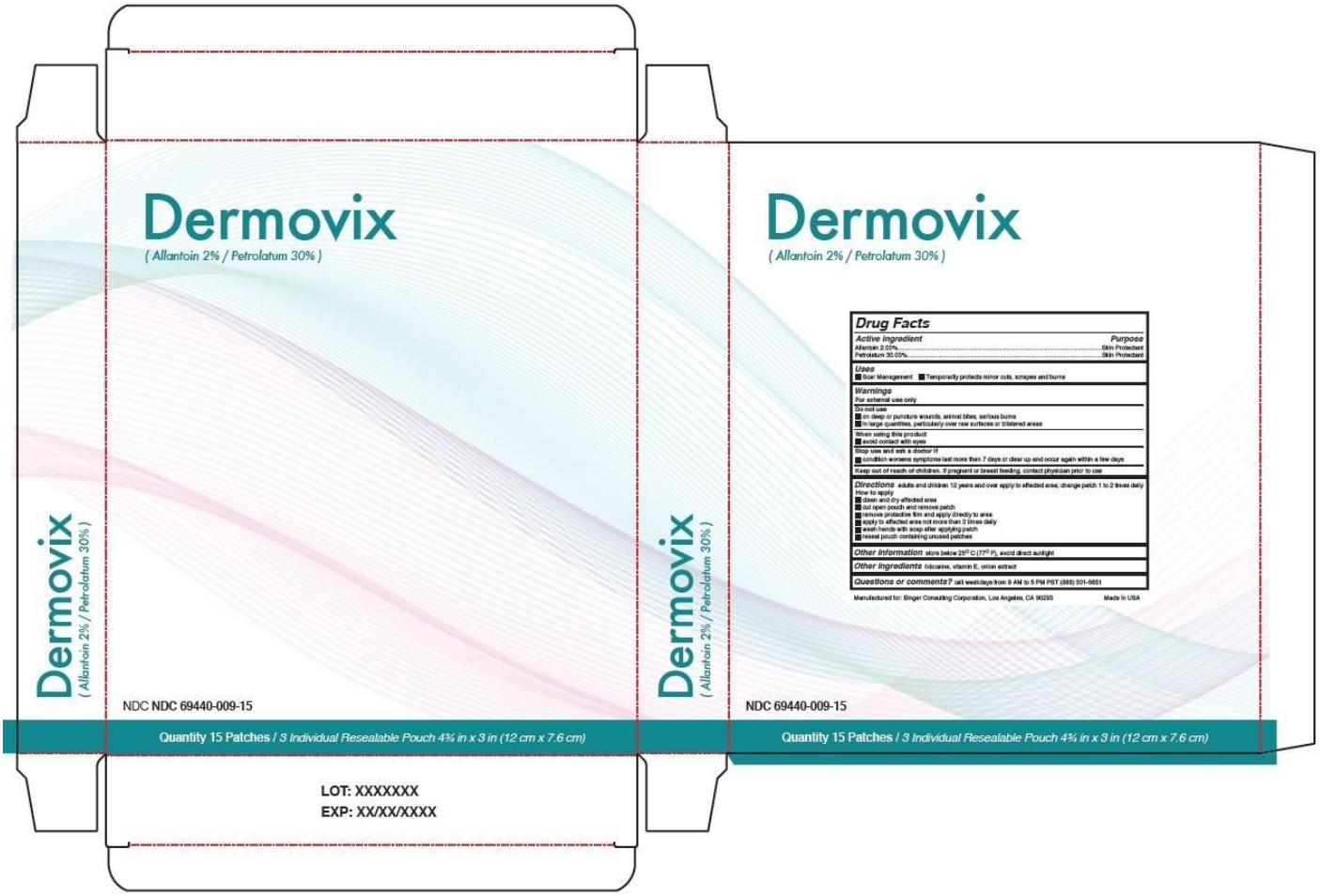
- clean and dry affected area
- cut open pouch and remove patch
- remove protective film and apply directly to area
- apply to affected area not more than 3 times daily
- wash hands with soap after applying patch
- reseal pouch containing unused patches

Other ingredients lidocaine, vitamin E, onion extract

Questions or comments? call weekdays from 9 AM to 5 PM PST (888) 501-5651

Other information store below 25°C (77° F), avoid direct sunlight

Packaging



DERMOVIX

allantoin,petrolatum patch

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69440-009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	2 g in 100 g
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	30 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
LIDOCAINE (UNII: 98PI200987)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
ONION (UNII: 492225Q21H)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69440-009-15	15 in 1 BOX	01/01/2015	
1		100 g in 1 PATCH; 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	01/01/2015	

Labeler - Binger Consulting Corporation (079635976)

Establishment

Name	Address	ID/FEI	Business Operations
Active Intelligence, LLC		080416593	manufacture(69440-009)

Revised: 10/2021 Binger Consulting Corporation