SKIN PROTECTANT PETROLATUM- petrolatum ointment Nationwide Pharmaceutical, 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.

SKIN PROTECTANT PETROLATUM OINTMENT

Drug Facts

Active ingredient

Petrolatum 44.28%

Purpose

Skin protectant

Uses

- Temporarily protects minor cuts, scrapes, and burns.
- Temporarily protects and helps relieve chapped or cracked skin and lips.
- Helps protect skin from drying effects of wind and cold weather.

Warnings

For external use only.

When using this product

do not get into eyes In case of contact, flush thoroughly with water.

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.

Do not use on

- deep or puncture wounds
- animal bites
- serious burns
- broken skin

Keep out of reach of children. In case of accidental ingestion contact a physician or Poison Control Center right away.

Directions

Apply as needed.

Other information

- Store at room temperature (59°-86°F)
- Do not use if inner seal is broken or missing

Inactive ingredients

cetostearyl alcohol, microcrystalline wax, mineral oil, and polysorbate 60

Questions or comments?

1-800-697-3329

PRINCIPAL DISPLAY PANEL - 454 GM Jar Label

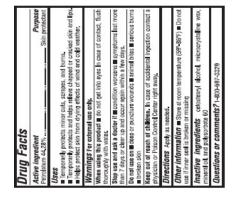
NDC 69375-0004-54

Nationwide Pharmaceutical

Skin Protectant Petrolatum Ointment

For dry, cracked or irritated skin

454 GM (16 OZ)





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454 GM (16 OZ)

Packaged For: Nationwide Pharmaceutical, LLC San Antonio, Tx 78232



SKIN PROTECTANT PETROLATUM

petrolatum ointment

Proc	luct	Info	rmation
	146		

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69375-004

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Petrolatum (UNII: 4T6H12BN9U) (Petrolatum - UNII:4T6H12BN9U)	Petrolatum	44.28 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
MICROCRYSTALLINE WAX (UNII: XOF597Q3KY)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	

Ш	Pa	ickaging				
#		Item Code	Package Description	Marketing Start Date	Marketing End Date	
		NDC:69375-004- 54	454 g in 1 JAR; Type 0: Not a Combination Product	03/01/2021		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH FINAL	part347	03/01/2021		

Labeler - Nationwide Pharmaceutical, LLC (079265801)

Revised: 2/2021 Nationwide Pharmaceutical, LLC