

PETROLEUM JELLY- petrolatum cream
Universal Distribution Center 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.

Nourishing Vitamin E Petroleum Jelly

Drug Facts

Active ingredient:

Petrolatum 30%

Purpose

Skin Protectant

Uses

- temporarily protects minor: • cuts • scrapes • burns
- temporarily protects and helps relieve chapped or cracked skin
- helps protect from the drying effects of wind and cold weather

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 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 right away.

Directions

apply as needed.

Inactive ingredients:

water (aqua), mineral oil, glycerin, cetearyl alcohol, cetyl alcohol, stearyl alcohol, glyceryl stearate, fragrance, polysorbate 80, laureth-20, BHT, methylparaben, propylparaben, dimethicone, tocopherol.

NEW

for Dry Skin

ABSORBS QUICKLY

Moisturizing

Replenish Skin Balance

Compare to the active ingredient of Vaseline® Petroleum Jelly Cream*

Helps restore skin's natural moisture

Distributed by:

Universal Distribution Center

96 Distribution Boulevard • Edison, NJ 08817

www.universaldc.com

Made in P.R.C

Packaging

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96 Distribution Boulevard Edison, NJ 08817
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ITEM#56134



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Made in P.R.C.

4.5 fl. oz. (133 ml)

PETROLEUM JELLY

petrolatum cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52000-123
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	30 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
MINERAL OIL (UNII: T5L8T28FGP)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
LAURETH-20 (UNII: 6SU0SC83AH)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
TOCOPHEROL (UNII: R0ZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-123-34	133 mL in 1 TUBE; Type 0: Not a Combination Product	07/27/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	07/27/2015	

Labeler - Universal Distribution Center LLC (019180459)

