

PETROLEUM- petrolatum jelly
Delon Laboratories (1990) Ltd

Petroleum Jelly

Active ingredient

White Petrolatum USP (100%)

Purpose

Skin protectant

Uses

- temporarily protects minor:
- cuts
- scrapes
- burns
- temporarily protects and helps relieve chapped or cracked skin and lips
- 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 or 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

Delon Petroleum Jelly USP 368g

DELON NDC 61734-040-03

Petroleum Jelly USP

Helps relieve chapped or cracked skin and lips

NET WT. 13 OZ.
(368 g)

SPACE FOR LOT # AND EXPIRY
DO NOT PRINT



0 59338 10203 0

14502-5

Drug Facts

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1583B-5

Made in Canada by: Delon Laboratories (1990) Inc
Pointe-Claire, QC, Canada H9R 1E2. www.labdalon.com

PETROLEUM

petrolatum jelly

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61734-040
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61734-040-03	368 g in 1 JAR; Type 0: Not a Combination Product	05/07/2010	11/30/2025
2	NDC:61734-040-04	212 g in 1 JAR; Type 0: Not a Combination Product	05/07/2010	08/31/2021
3	NDC:61734-040-05	113 g in 1 JAR; Type 0: Not a Combination Product	10/04/2014	12/04/2014
4	NDC:61734-040-06	10 g in 1 TUBE; Type 0: Not a Combination Product	10/04/2014	12/04/2014
5	NDC:61734-040-07	100 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/07/2010	11/08/2017
6	NDC:61734-040-08	105 g in 1 JAR; Type 0: Not a Combination Product	10/04/2014	12/04/2014
7	NDC:61734-040-09	255 g in 1 JAR; Type 0: Not a Combination Product	10/04/2014	12/04/2014

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	05/07/2010	11/30/2025

Labeler - Delon Laboratories (1990) Ltd (248364184)

Establishment

Name	Address	ID/FEI	Business Operations
Laboratoires Delon		208896216	pack(61734-040) , manufacture(61734-040) , label(61734-040)

Revised: 10/2023

Delon Laboratories (1990) Ltd