

BABYFRESH PETROLEUM JELLY- petrolatum jelly
Delon Laboratories (1990) Ltd

Baby Fresh Petroleum Jelly

Active Ingredient

White Petrolatum USP (99.7%)

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

Inactive ingredient

fragrance

Delon Baby Fresh Petroleum Jelly 13 oz (368 g)



Drug Facts	
Active ingredient White Petrolatum USP (99.7%)	Purpose Skin protectant
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Directions apply as needed	
Inactive ingredient fragrance	

MADE IN CANADA
EBOLENE PRODUCTS CO.
HARTFORD, CT 06114
U.S.A.
18 W.P.B.C.
Wrenley HAG OAS
England
Made in Canada
14483B-4

Eboline Baby Fresh Petroleum Jelly 13 oz (368 g)



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Distributed by:
Eboline Products Co.
Hartford, CT 06114
U.S.A.
18 W.P.B.C.
Wrenley HAG OAS
England
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BABYFRESH PETROLEUM JELLY

petrolatum jelly

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61734-050
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)			PETROLATUM	99.7 g in 100 g
Inactive Ingredients				
Ingredient Name			Strength	
TALC (UNII: 7SEV7J4R1U)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61734-050-03	368 g in 1 JAR; Type 0: Not a Combination Product	05/07/2010	05/31/2024
2	NDC:61734-050-04	212 g in 1 JAR; Type 0: Not a Combination Product	05/07/2010	07/31/2023
3	NDC:61734-050-05	90 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/04/2014	12/04/2014
4	NDC:61734-050-06	170 g in 1 JAR; Type 0: Not a Combination Product	02/14/2019	11/30/2020
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M016	05/07/2010	05/31/2024

Labeler - Delon Laboratories (1990) Ltd (248364184)

Establishment			
Name	Address	ID/FEI	Business Operations
Laboratoires Delon		208896216	label(61734-050) , pack(61734-050) , manufacture(61734-050)

Revised: 10/2023

Delon Laboratories (1990) Ltd