

PRINCESS PETROLEUM- white petrolatum jelly
Ultra Distributors Inc.

Princess Petroleum Jelly

Active Ingredient

White petrolatum USP (100%)

Purpose

Skin protectant

Uses

helps protect chapped skin

Warnings

For external use only

When using this product

- do not get into eyes

See a doctor if condition lasts more than 7 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

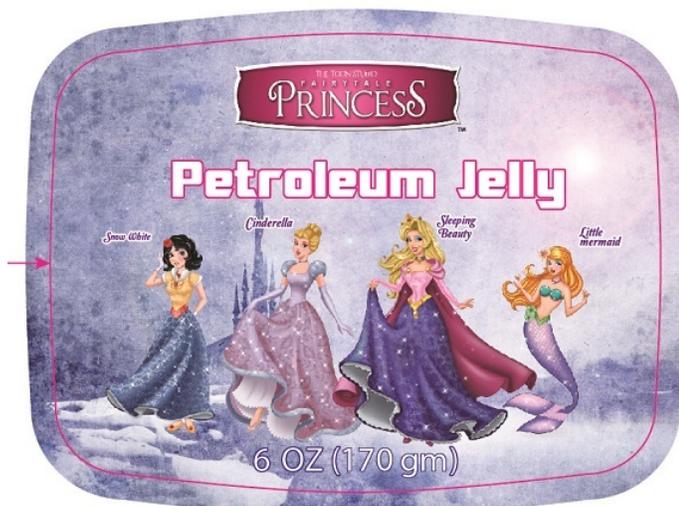
Inactive ingredients

None

PRINCIPAL DISPLAY PANEL

PRINCESS PETROLEUM JELLY

NET WT 6 OZ (170 gm)



Distributed by: Ultra Distributors Inc.
 Somerset NJ 08873, USA
 License by The Walt Disney Company
 Snow White, Cinderella and Sleeping Beauty
 based on Jr. FFBrooks and Record Guild of America.
 Little Mermaid based on Hans Christian Andersen.
 All rights reserved © 2019 United Trademark Holding

Item No. 4643
 7 88268 184543 7

Drug Facts	
Active ingredient	Purpose
White petrolatum USP (100%).....	Skin protectant
Use helps protect chapped skin	
Warnings	
For external use only	
When using this product • do not get into eyes	
See a doctor if condition lasts more than 7 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	

Made in India DNHC-92

PRINCESS PETROLEUM

white petrolatum jelly

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78495-004
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:78495-004-01	113 g in 1 JAR; Type 0: Not a Combination Product	12/01/2020	
2	NDC:78495-004-02	170 g in 1 JAR; Type 0: Not a Combination Product	12/01/2020	
3	NDC:78495-004-03	226 g in 1 JAR; Type 0: Not a Combination Product	12/01/2020	
4	NDC:78495-004-04	368 g in 1 JAR; Type 0: Not a Combination Product	12/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	12/01/2020	

Labeler - Ultra Distributors Inc. (007160073)

Registrant - Jell Pharmaceuticals Pvt. Ltd. (726025211)

Establishment

Name	Address	ID/FEI	Business Operations
Jell Pharmaceuticals Pvt. Ltd.		726025211	manufacture(78495-004)

Revised: 12/2024

Ultra Distributors Inc.