

**PETROLEUM JELLY- white petrolatum jelly**  
**Aldermed Inc.**

-----  
**Petroleum Jelly**

***Active ingredient***

White petrolatum USP 100%

***Purpose***

Skin protectant

***Uses***

temporarily protects minor

- cuts
- scrapes
- burns
- temporarily protects and chafed, chapped or cracked skin
- helps prevent against the drying effects of wind and cold weather

***Warnings***

**For external use only**

**Do not use on**

- deep or puncture wounds
- animal bites
- serious burns

**When using this product**

- do not get into eyes

**Stop use and ask a doctor if**

- condition worsens
- symptoms last for more than 7 days or clear up and occur again within a few days

**Keep out of reach of children.**

In swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

***Directions***

apply as needed

## Other information

- store at 15-30 C (59-86F)

## Inactive ingredients

none

## Questions or comments?

Call 1-833-605-84-74 9am-5 pm EST M-F

## Distributor

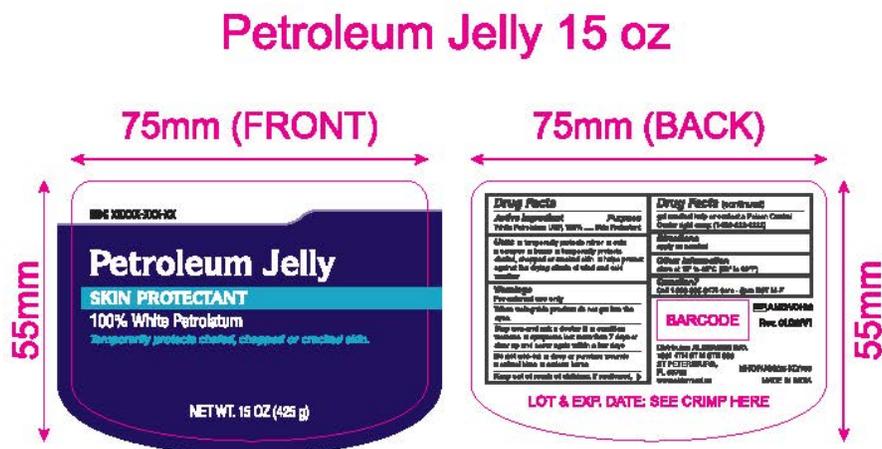
Distributor:

ALDERMED INC.

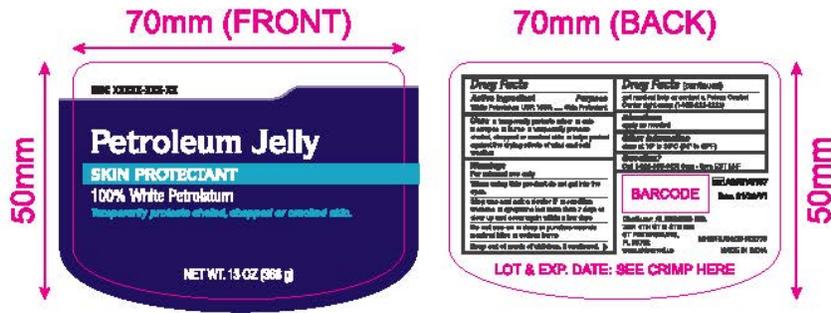
7901 4TH ST N STE 300 ST PETERSBURG, FL 33702

www.aldermed.us MADE IN INDIA MH/DRUGS/25-KD/739

## Package Label



# Petroleum Jelly 13 oz



## PETROLEUM JELLY

white petrolatum jelly

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:87236-009
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>WHITE PETROLATUM</b> (UNII: B6E5W8RQJ4) (WHITE PETROLATUM - UNII:B6E5W8RQJ4)	WHITE PETROLATUM	1 g in 1 g

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87236-009-02	56.6 g in 1 TUBE; Type 0: Not a Combination Product	12/01/2025	
2	NDC:87236-009-13	368 g in 1 JAR; Type 0: Not a Combination Product	12/01/2025	
3	NDC:87236-009-15	425 g in 1 JAR; Type 0: Not a Combination Product	12/01/2025	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	12/01/2025	

**Labeler** - Aldermed Inc. (144887712)

**Registrant** - Aldermed Inc. (144887712)

Revised: 2/2026

Aldermed Inc.