# SKIN-LUBE- alcohol ointment Performance Health, 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.

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#### **Skin-Lube Ointment**

## **Drug Facts**

## **Active Ingredients**

Petrolatum 85%

Dimethicone 1.5%

## **Purpose**

Skin Protectant

#### Uses

Temporarily protects minor - cuts - scrapes - burns -- Helps prevent, temporarily protects, and helps relieve chafed, chapped or cracked skin. Helps prevent and protect from 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 more than 7 days or clear up and occur again within few days.

#### **Directions:**

Apply as needed.

# Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

#### Other Information:

Store in a cool dry place with lid closed tightly

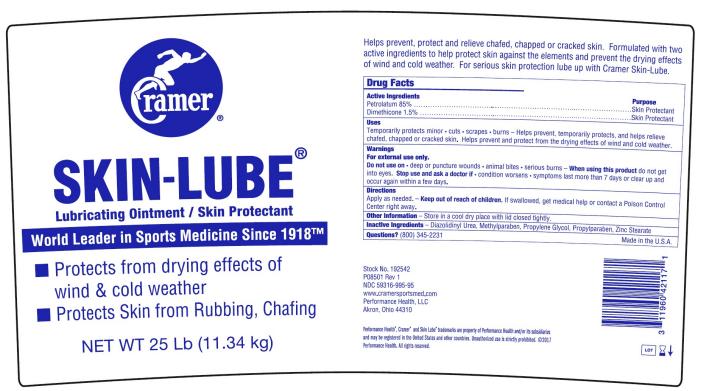
#### **Inactive Ingredients**

Diazolidinyl Urea, Methylparaben, Propylene Glycol, Propylparaben, Zinc Stearate

#### Questions?

(800) 345-2231

**Package Labeling: 59316-995-95** 



# SKIN-LUBE alcohol ointment Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:59316-995 Route of Administration TOPICAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength

PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	850 mg in 1 g
DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10)	DIMETHICONE	15 mg in 1 g

Inactive Ingredients				
Ingredient Name	Strength			
METHYLPARABEN (UNII: A218C7H19T)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
PROPYLPARABEN (UNII: Z8IX2SC10H)				
ZINC STEARATE (UNII: H92E6QA4FV)				
DIAZOLIDINYL UREA (UNII: H5RIZ 3MPW4)				

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:59316- 995-21	78 g in 1 JAR; Type 0: Not a Combination Product	05/18/2017	12/31/2024			
2	NDC:59316- 995-95	11340 g in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2017	12/31/2024			
3	NDC:59316- 995-96	454 g in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2017	12/31/2024			
4	NDC:59316- 995-97	2270 g in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2017	12/31/2024			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part347	05/18/2017	12/31/2024		

# Labeler - Performance Health, LLC (794324061)

Revised: 8/2022 Performance Health, LLC