PROSKI ZINCO SKIN PROTECTANT- zinc oxide ointment PHARMAMED USA INC

Zinc Oxide Ointment

ACTIVE INGREDIENT

Zinc Oxide 20%

PURPOSE

Skin Protectant

USE

Helps treat chafed skin associated with diaper rash

Dries the oozing and weeping of poison: • ivy • oak •sumac

WARNINGS

For External Use Only.

When using this product

Avoid contact with 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

Keep out of reach of children

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

DIRECTIONS

- For diaper rash: Change wet and soiled diapers promptly, cleanse the diaper area, and allow to dry.
- Apply ointment liberally as often as necessary with each diaper change, especially at bedtime or anytime when exposure to wet diapers may be prolonged.
- For poison ivy, oak, and sumac: Apply as needed.

OTHER INFORMATION

- Store at room temperature between 15° 30°C (59° 86°F)
- Avoid excessive heat
- Tamper evident. Do not use if seal is damaged.

INACTIVE INGREDIENTS

CETETH-20, CETOSTEARYL ALCOHOL, LIGHT MINERAL OIL, WHITE PETROLATUM

QUESTIONS OR COMMENTS?

(754)-200-8994 (Mon-Fri 8 AM to 4 PM EST)

PACKAGE/LABEL PRINCIPAL DISPLAY PANEL

NDC: 84289-119-20

ZINC OXIDE OINTMENT

Skin Protectant

Net Wt 1 oz. (28.4 g)

GOA/DRUGS/571





PROSKI ZINCO SKIN PROTECTANT

zinc oxide ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84289-119

Route of Administration TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	200 mg in 1 g		

Inactive Ingredients			
Ingredient Name	Strength		
CETETH-20 (UNII: 1835H2IHHX)			
LIGHT MINERAL OIL (UNII: N6K5787QVP)			
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)			
WHITE PETROLATUM (UNII: B6E5W8RQJ4)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:84289-119- 20	1 in 1 CARTON	04/01/2025			
1		28.4 g in 1 TUBE; Type 0: Not a Combination Product				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M016	04/01/2025		

Labeler - PHARMAMED USA INC (065607328)

Revised: 3/2025 PHARMAMED USA INC