

ZINC OXIDE 20%- zinc oxide ointment
Pharmaceutica North America, Inc.

Active ingredient

Zinc Oxide 20% w/w

Purpose

Skin Protectant

Uses

Dries the oozing and weeping of poison: • Ivy • Oak • Sumac

Warnings

For External Use Only

When using this product

- Do not get into eyes
- Do not use over large areas of the body

Stop use and ask a doctor if

- The condition worsens
- Symptoms last more than 7 days or clear up and occur again within a few days
- If you are allergic to any of these ingredients

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 poison ivy, oak, and sumac: apply as needed.

Other information

- Store at room temperature between 15°-30°C (59°-86°F)
- Avoid excessive heat

Inactive Ingredients

Cetomacrogol 1000, Cetostearyl Alcohol, Light Mineral Oil, White Soft Paraffin (Petrolatum)

Questions or comments? (888)788-6472

Product label





NDC: 45861-0082-06

Zinc Oxide Ointment

20%



NET WT 2.11 OZ (60g)

Drug Facts

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ZINC OXIDE 20%

zinc oxide ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:45861-082
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	20 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-20 (UNII: I835H2IHHX)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:45861-082-06	1 in 1 BOX	02/01/2025	

1	60 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	02/01/2025	

Labeler - Pharmaceutica North America, Inc. (962739699)

Revised: 3/2025

Pharmaceutica North America, Inc.