ZINC OXIDE 20%- zinc oxide 20% ointment
SOLA Pharmaceuticals

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.

Zinc Oxide 20% Ointment

Active Ingredient

Zinc Oxide 20%

Purpose

Skin protectant

Uses

Skin Protectant
- Helps treat and prevent diaper rash
- Dries the ozzing 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

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

Other Information
- Store at room temperature
- Avoid excessive heat and humidity

Inactive ingredients

Cetomacrogol 1000, Cetostearyl alcohol, Mineral oil, Petrolatum

Questions of comments? Call 1-866-747-7365

Manufactured for : SOLA Pharmaceuticals

Baton Rouge, LA 70810

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Zinc Oxide 20% Ointment

NDC 70512-103-30

Qty: 28.4g

ZINC OXIDE 20%

zinc oxide 20% ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70512-103
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: I835H2IHHX)	
PETROLATUM (UNII: 4T6H12BN9U)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
MINERAL OIL (UNII: T5L8T28FGP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70512-103-30	1 in 1 CARTON	04/22/2021	
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 final	part347	04/22/2021	

Labeler - SOLA Pharmaceuticals (080121345)

Revised: 7/2022 SOLA Pharmaceuticals