ZINC OXIDE 20%- zinc oxide ointment
Pharmacy Value Alliance, 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.

Zinc Oxide Ointment 20%

Drug Facts

Active Ingredient

Zinc Oxide 20%

Purpose

Skin Protectant

Uses

- Helps treat and prevent diaper rash
- Protects chafed skin due to diaper rash - helps protect skin from wetness
- Protects and dries the oozing and weeping of poison ivy, poison oak and poison sumac
- Helps prevent and temporarily protect chafed, chapped, cracked or windburned skin and lips

Keep out of the reach of children

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

Warnings

For External Use Only

- When using this product avoid contact with eyes
- Stop use and ask a doctor if condition worsens or does not improve within 7 days. This may be a sign of a serious condition.

Directions

- 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 betime or anytime when exposure to wet diapers may be prolonged

Other Information

- Store at controlled room temperature 20° - 25°C (68° - 77°F)

Inactive Ingredients

Mineral Oil, Petrolatum

Distributed By:

Pharmacy Value Alliance, LLC.

407 East Lancaster Avenue, Wayne, PA. 19087

www.emersongroup.com

Product of PRC

Packaging

OUTER BOX

INNER TUBE

ZINC OXIDE 20%

zinc oxide ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-159
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.2 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-159-02	1 in 1 BOX	07/19/2016	
1		57 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	07/19/2016	

Labeler - Pharmacy Value Alliance, LLC (101668460)

Registrant - Trifecta Pharmaceuticals USA LLC (079424163)

Revised: 5/2023 Pharmacy Value Alliance, LLC