ZINC OXIDE 20%- zinc oxide ointment
Chain Drug Marketing Association Inc.

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.

Quality Choice Zinc Oxide Ointment USP

Drug Facts

Active Ingredient

Zinc Oxide 20%

Purpose

Skin Protectant

Uses

- Protects skin
- Protects and releives chafed skin due to diaper rash - helps seal out wetness
- Dries the oozing and weeping of poison ivy, oak and sumac

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

- Apply ointment liberally as often as necessary
- For the treatment of diaper rash, change wet and soiled diapers promptly
- Cleanse the diaper area and allow to dry
- Apply with each diaper change, especially at bedtime or anytime when exposure to wet diaper may be prolonged

Other Information

- Store at controlled room temperature 15° - 30°C (59° - 86°F)

Inactive Ingredients

Mineral Oil, Petrolatum

Distributed By:

Distributed by CDMA Inc.

43157 W 9 Mile Rd

Novi, MI. 48375

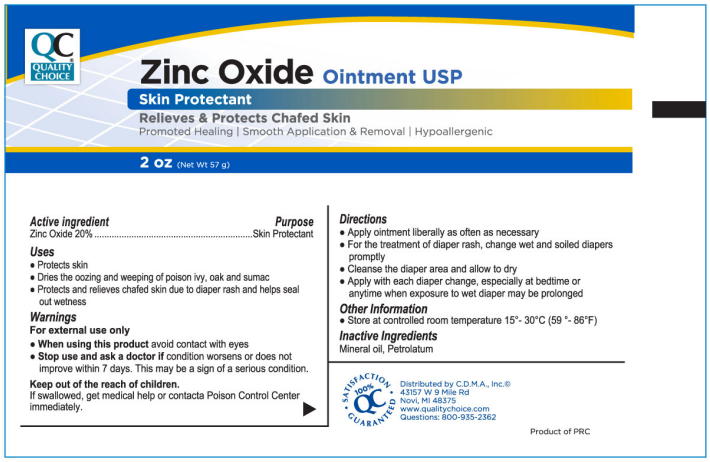
Product of PRC

www.qualitychoice.com

Questions: 800-935-2362

OUTSIDE BOX

INNER TUBE



ZINC OXIDE 20%

zinc oxide ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-686
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:63868-686-02	1 in 1 BOX	08/14/2020	
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	08/14/2020	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Registrant - Trifecta Pharmaceuticals USA LLC (079424163)

Revised: 5/2023 Chain Drug Marketing Association Inc.