MEDPURA ZINC OXIDE 20%- medpura zinc oxide 20% ointment Akron Pharma 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.

MEDPURA
Zinc Oxide 20% Ointment
Regular Strength

Active Ingredient

Zinc Oxide 20%

Purpose

Skin Protectant

Uses

- helps treat and prevent diaper rash
- protects chafed skin due to diaper rash and helps seal out wetness
- protects and dries the oozing and weeping of poison ivy, poison oak, and poison sumac.

Warnings

For External Use Only

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

if condition worsens or does not improve within 7 days. This may be a sign of a serious condition.

Keep out of reach of children

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

Directions

- change wet or soiled diapers promptly
- cleanse the diaper area and allow to dry
- apply ointment liberally, as often as necessary, with each diaper change and

especially at bedtime or anytime when exposure to wet diapers may be prolonged.

Other Information

- store at room temperature 20°-25°C (68°-77°F)
- may stain clothing

Inactive Ingredients

alovera gel, mineral oil, white petrolatum

Questions and Comments?

Call toll-free 1-877-225-6999







MEDPURA ZINC OXIDE 20%

medpura zinc oxide 20% ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71399-0245
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	20 g in 100 g		

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
MINERAL OIL (UNII: T5L8T28FGP)		
WHITE PETROLATUM (UNII: B6E5W8RQJ4)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:71399- 0245-2	1 in 1 CARTON	09/12/2022		
1		56.7 g in 1 TUBE; Type 0: Not a Combination Product			
2	NDC:71399- 0245-3	1 in 1 CARTON	11/04/2021		
2		85 g in 1 TUBE; Type 0: Not a Combination Product			
3	NDC:71399- 0245-6	454 g in 1 JAR; Type 0: Not a Combination Product	11/04/2021		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part347	11/04/2021		

Labeler - Akron Pharma Inc. (067878881)

Revised: 3/2023 Akron Pharma Inc.