MAXIMUM STRENGTH DIAPER RASH- zinc oxide ointment ointment CVS Pharmacy

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.

Active ingredient

Zinc Oxide, 40%

Purpose

Skin protectant

Uses

helps treat and prevent diaper rash, Protects chafed skin due to diaper rash and helps seal out wetness

Warnings

For external use only.

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

condition worsens or does not improve after 7 days

Keep out of reach of children

to prevent accidential ingestion. If swallowed, seek medical help or call Poison Control Center immediately.

Directions for Use

change wet and soiled diaper immediately, cleanse the diaper area and allow to dry, apply ointment liberally as often as necessary with each diaper change and especially when exposured to wet diapers for a prolonged period of time, such as bedtime

Other information

Storage temperature: not to exceed 30oC (86oF),

Remove foil seal from the tube's tip,

Use with infants, children, and adults,

Will stain clothing and fabric

Inactive ingredients

Package label





MAXIMUM STRENGTH DIAPER RASH

zinc oxide ointment ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-849
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	40 g in 100 g

Inactive Ingredients			
Ingredient Name	Strength		
PETROLATUM (UNII: 4T6H12BN9U)	31 g in 100 g		
MINERAL OIL (UNII: T5L8T28FGP)	20 g in 100 g		
PARAFFIN (UNII: 19 O 0 E 3 H 2 Z E)	1 g in 100 g		
CASTOR OIL (UNII: D5340 Y2I9 G)	3.5 g in 100 g		
BALSAM PERU (UNII: 8 P5F8 8 10 CY)	4.5 g in 100 g		

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Pa	ckaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1 N	IDC:59779-849-04	113 g in 1 TUBE; Type 0: Not a Combination Product	07/22/2016	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	07/22/2016	

Labeler - CVS Pharmacy (062312574)

Establishment			
Name	Address	ID/FEI	Business Operations
Unipack, Inc.		009248480	manufacture(59779-849)

Revised: 1/2021 CVS Pharmacy