DIAPER RASH- zinc oxide paste H E B

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.

Drug Facts 020.002 020AC/AD

Active Ingredients

Zinc oxide 40%

Purpose

Skin protectant

use

- 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

• do not get into 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 or contact a Poison Control Center right away.

Directions

- change wet and soilded diapers promptly
- cleanse the diaper area
- allow to dry
- apply ointment liberally as often as necessary, with each diaper change, especially at bedtime or anytime when exposure to wet diapers may be prolonged

Other information

store between 200 and 250C (680 and 770F)

Inactive ingredients

petrolatum, cod liver oil, lanolin, Zea mays (corn) starch, glycerin, sorbitan sesquioleate, beeswax,

tocopheryl acetate, fragrance

Adverse Reactions

DISTRIBUTED BY: H-E-B, SAN ANTONIO, TN 78204

LOT NUMBER ON PACKAGE

QUESTIONS? 1-888-593-0593

MADE IN USA WITH US AND IMPORTED PARTS.

WE HOPE YOU ARE SATISFIED WITH

THIS PRODUCT. IF NOT, WE WILL

CHEERFULLY REFUND YOUR MONEY

KEEP THIS CARTON FOR FULL PRODUCT INFORMATION

PRINCIPAL DISPLAY PANEL

HILL COUNTRY

ESSENTIALS

Diaper

Rash Paster

SKIN PROTECTANT

40% ZINC OXIDE

Helps Soothe &

Prevent Diaper Rash

Pomada para

Rozaduras

PROTECTOR DE LA PIEL

CON OXIDO DE

ZINCE 40%

FREE FROM

parapbens

phthalates

dyes

NET WT. 4 OZ

CONT. NET. 113 g



Diaper Rash Paste

SKIN PROTECTANT

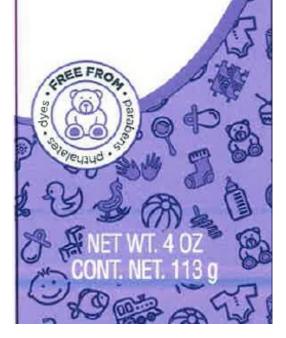
40% ZINC OXIDE

Helps Soothe & Prevent Diaper Rash

Pomada para Rozaduras

PROTECTOR DE LA PIEL

CON OXIDO DE ZINC 40%



DIAPER RASH

zinc oxide paste

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	532 mg in 1 g	

Inactive Ingredients				
Ingredient Name	Strength			
PETROLATUM (UNII: 4T6H12BN9U)				
COD LIVER OIL (UNII: BBL281NWFG)				
LANOLIN (UNII: 7EV65EAW6H)				
STARCH, CORN (UNII: O8232NY3SJ)				
GLYCERIN (UNII: PDC6A3C0OX)				
SORBITAN SESQUIOLEATE (UNII: 0 W8 RRI5W5A)				
YELLOW WAX (UNII: 2ZA36H0S2V)				
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)				

1	Packaging			
#	# Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-020-26	1 in 1 CARTON	09/10/2010	
1		113 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	09/10/2010	

Labeler - HEB (007924756)

Registrant - Vi-Jon (790752542)

Establishment				
Name	Address	ID/FEI	Business Operations	
Vi-jo n		790752542	manufacture(37808-020)	

Establishment				
Name	Address	ID/FEI	Business Operations	
Memphis Contract Packaging		185390010	manufacture(37808-020)	

Revised: 4/2020 HE B