MY FAIR BABY DIAPER RASH- zinc oxide cream Anicare Pharmaceuticals Pvt. Ltd.

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.

diaper rash cream

Active Ingredient

Zinc Oxide 13%

Purpose

Skin Protection

Use

- Helps treat and prevent diaper rash
- Soothes and relieves irritated skin
- Protects your baby's skin from 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 within 7 days or symptoms 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 soiled diapers promptly
- Cleanse the diaper area
- Allow to dry
- Ointment should be applied at every diaper change, specially at bedtime or anytime when exposure to wet diapers may be prolonged

Other Information

Store at 20° to 25°C (68° to 77°F)

Inactive Ingredients

Water, Mineral Oil, Petrolatum, Cetyl Alcohol, Ceteareth-25, Glyceryl Monostearate, Beeswax, Microcrystaline Wax, Fragrance, Dimethicone, Aloe Barbadensis Leaf Extract (Aloe Vera), Tocopheryl Acetate, Methylparaben, Propylparaben, Methylchloroisothiazolinone, Methylisothiazolinone, Carbomer

Carton Label





Diaper Rash Cream

THE RIGHT CHOICE OF MOMS

0321/01(45)

Instant Relief

Drug Facts

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Drug Facts (continued)

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Zinc Oxide

ITEM NO: 10321





Specially Manufactured for **DELTA BRANDS INC.** Larchmont, NY 10538 www.deltabrands.com MADE IN CHINA *This product is not manufactured or distributed by Johnson & Johnson

MY FAIR BABY DIAPER RASH

zinc oxide cream

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47046-179	
Route of Administration	TOPICAL			

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

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
MINERAL OIL (UNII: T5L8T28FGP)		
PETROLATUM (UNII: 4T6H12BN9U)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
CETEARETH-25 (UNII: 8FA93U5T67)		
GLYCERYL MONOSTEARATE (UNII: 230 OU9 XXE4)		
WHITE WAX (UNII: 7G1J5DA97F)		
MICRO CRYSTALLINE WAX (UNII: XOF597Q3KY)		
DIMETHICO NE (UNII: 92RU3N3Y1O)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
PROPYLPARABEN (UNII: Z8 IX2SC1OH)		
METHYLCHLORO ISO THIAZO LINO NE (UNII: DEL7T5QRPN)		
METHYLISO THIAZO LINO NE (UNII: 229 D0 E1QFA)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:47046-179-02	1 in 1 CARTON	08/30/2011		
1	NDC:47046-179-01	28 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	08/30/2011		

Labeler - Anicare Pharmaceuticals Pvt. Ltd. (916837425)

Revised: 12/2020 Anicare Pharmaceuticals Pvt. Ltd.