OLP DIAPER RASH CREAM A AND D- zinc oxide cream Ohio Lab Pharma

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 Ingredients

Zinc Oxide 10%

Stop using this product and ask a doctor

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days

Uses

- helps treat and prevent diaper rash
- protects minor skin irritation due to diaper rash and helps seal out wetness

Warning

- For external use only
- when using this product, do not get it into the eye

questions

www.ohiolabpharma.us

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, and allow to dry
- apply cream liberally as often as necessary, with each diaper change, especially at bedtime or anytime when exposure to wet diapers may be prolonged

store between 20°C to 25 °C (68° to 77°F)

Inactive Ingredients

cod liver oil (contains vitamin A & Vitamin D), propylene glycol, vitamin E, methylparaben, propylparaben, stearic acid, trolamine, ceteareth 20, cetostearyl alcohol, EDTA, mineral oil

purpose

diaper rash cream Maximum Strength NET WEIGHT 0.7 oz (20g)

NDC#70648-444-01



Product Information							
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC		NDC:70648	C:70648-444		
Route of Administration	TOPICAL						
Active Ingredient/Active Mo	oiety						
Ing		Basis of Streng	gth S	Strength			
ZINC OXIDE (UNII: SOI2LOH54Z) (Z	INC CATION - UNII:13S1S8SF37)		ZINC CATION	10 g	in 100 g		
	Ingredient Name			St	rength		
Inactive Ingredients	Ingradiant Nama			St	rangth		
.ALPHATOCOPHEROL (UNII: H4N					0		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)							
COD LIVER OIL (UNII: BBL281NWF	G)						
CETOSTEARYL ALCOHOL (UNII: 2	DMT128M1S)						
TROLAMINE (UNII: 903K93S3TK)							
MINERAL OIL (UNII: T5L8T28FGP)							
	C86K)						
EDETATE DISODIUM (UNII: 7FLD91							
EDETATE DISODIUM (UNII: 7FLD9 1 STEARIC ACID (UNII: 4ELV7Z65AP)							

Product Characteristics								
Color		white	Score					
Shape			Size					
Flavor			Imprint Code					
Contains								
Packaging								
# Item Code	I	Package Description		Marketing Start Date	Marketing End Date			
1 NDC:70648-444-01	1 in 1 CARTON		10/12/2017					
1	20 g in 1 TUBE; Type 0: Not a Combination Product							
Marketing Information								
Marketing Category	Applicatio	n Number or Monog	raph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part347			10/12/2017				

Labeler - Ohio Lab Pharma (080215854)

Establishment

Name	Address	ID/FEI	Business Operations
Ohio Lab Pharma		080215854	manufacture(70648-444)

Revised: 11/2018

Ohio Lab Pharma