DIAPER RASH- zinc oxide cream
Target Corporation

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.

Up & Up 226.002 226AD

Active ingredient

Zinc oxide 13%

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 soiled diaper promplty
- 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 20º and 25ºC (68º and 77º F)

Inactive ingredients

water, mineral oil, petrolatum, beeswax, dimethicone, sorbitan sesquioleate, microcrystalline wax, PEG-30 dipolyhydroxystearate, Aloe barbadensis leaf juice, glycerin, tropolone, tocopheryl acetate, 1,2-hexanediol, caprylyl glycol, magnesium sulfate, potassium hydroxide, phenoxyethanol

Questions

Call 1-800-910-6874

Disclaimer

This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company, distributor of Destine Diaper Rash Cream

Adverse Reactions

Distributed b Target Corporation

Minneapolis, MN 55403

Made in the U.S.A. with U.S.A. and foreign components

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principal display panel

NDC 11673-226-26

Compare to Destin

Diaper Rash Cream

fast relief

zinc oxide

creamy

diaper rash

ointment

skin protectant

helps soothe, heal &

prevent diaper rash

dermatologisted tested

hypoallengenic

fragrance free

up & up

NET WT 4 OZ (113 g)

DIAPER RASH

zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-226
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	130 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
MICROCRYSTALLINE WAX (UNII: XOF597Q3KY)	
PEG-30 DIPOLYHYDROXYSTEARATE (UNII: 9713Q0S7FO)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
TROPOLONE (UNII: 7L6DL16P1T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
MAGNESIUM SULFATE MONOHYDRATE (UNII: E2L2TK027P)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-226-26	1 in 1 CARTON	08/08/1994	
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	08/08/1994	

Labeler - Target Corporation (006961700)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(11673-226)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(11673-226)