DIAPER RASH- zinc oxide paste
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.

Diaper Rash Ointment 40% Zinc Oxide
020.002/020AC-AD

Active ingredient

Zinc oxide 40%

Purpose

Skin Protectant

Uses

• helps treat and prevent diaper rash
• protects chafed skin due to the 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 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 20º and 25ºC (68º and 77ºF)

Inactive ingredients

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

Questions

Call 1-800-910-6874

Claims

This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company, distributor of Destine Maximum Strength

40% Zinc Oxide Diaper Rash Paste.

Adverse Reactions

Distributed by Target Corporation

Minneapolis, MN 55403

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

Principal display panel

NDC 11673-020-26

Compare to Desitin

maximum strength

diaper rash paste*

40% zinc oxide

skin protectant

pediatrician tested

seals out wetness

hypoallergenic

paraben free

up & up

NET WT 4 OZ (113.3 g)

DIAPER RASH

zinc oxide paste

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	400 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: 0W8RRI5W5A)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-020-26	1 in 1 PACKAGE	03/24/2008	
1		113.3 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	03/24/2008	

Labeler - Target Corporation (006961700)

Registrant - Vi Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		790752542	manufacture(11673-020)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, LLC		088520668	manufacture(11673-020)

Revised: 4/2023 Target Corporation