

**QUALITY CHOICE DIAPER RASH- zinc oxide ointment**  
**Chain Drug Marketing Association Inc.**

*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.*

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**Quality Choice Diaper Rash Ointment**

**Drug Facts**

**Active Ingredient**

Zinc Oxide 40%

**Purpose**

Skin Protectant

**Uses**

- Helps treat and prevent diaper rash
- Protects chafed skin due to diaper rash and helps protect skin from wetness

**Keep out of the reach of children.**

If swallowed, get medical help or contact a Poison Control Center immediately.

**Warnings**

**For External Use Only**

When using this product

- Avoid contact with eyes

This product contains trace amounts of naturally occurring lead.

**Directions**

- Change wet and soiled diapers promptly
- Cleanse the diaper area and 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.

**Stop use and ask a doctor if**

- Condition worsens

- Symptoms last more than 7 days or clear up and occur again with a few days

**Other Information:**

- Store at controlled room temperature 20° - 25°C (68° - 77°F)

**Questions**

1-800-935-2362

**Inactive Ingredients**

BHA, Cod liver oil (high in vitamin A and D), Frangrance, Lanolin, Mineral Oil, Methylparaben, Petrolatum, Purified water, Talc, Tartrazine yellow #4 (FD and C yellow #4)

**Helps treat and prevent diaper rash. Protects skin. Relieves Chafing. Promotes Healing**

Distributed by:

C.D.M.A., Inc. ©

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www.qualitychoice.com

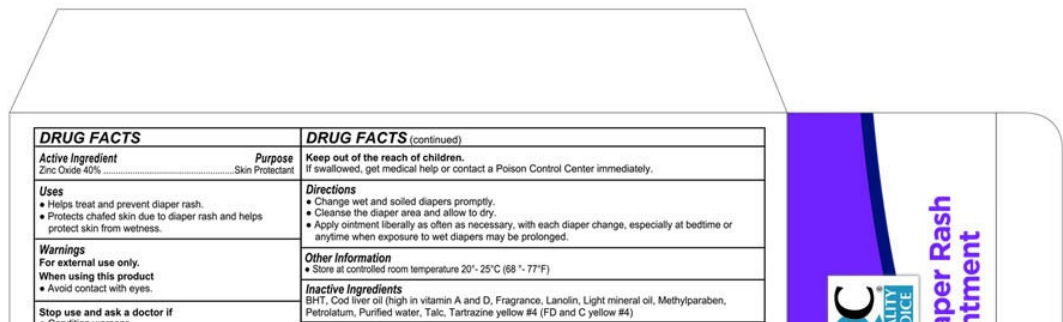
Product of PRC

**Other Information**

This product is not manufactured or distributed by Johnson & Johnson Consumer Company Inc. owner of the registered trademark Desitin®.

**Packaging**

OUTER BOX





INNER TUBE



**QUALITY CHOICE DIAPER RASH**

zinc oxide ointment

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63868-948
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	40 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
TALC (UNII: 7SEV7J4R1U)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
LANOLIN (UNII: 7EV65EAW6H)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
COD LIVER OIL (UNII: BBL281NWFG)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PETROLATUM (UNII: 4T6H12BN9U)	
WATER (UNII: 059QF0KO0R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-948-04	113 g in 1 TUBE; Type 0: Not a Combination Product	12/19/2019	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	12/19/2019	

**Labeler** - Chain Drug Marketing Association Inc. (011920774)**Registrant** - Trifecta Pharmaceuticals USA LLC (079424163)

Revised: 5/2023

Chain Drug Marketing Association Inc.