

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

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), Fragrance, 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:

CDMA Inc.

Novi, MI. 48375 USA

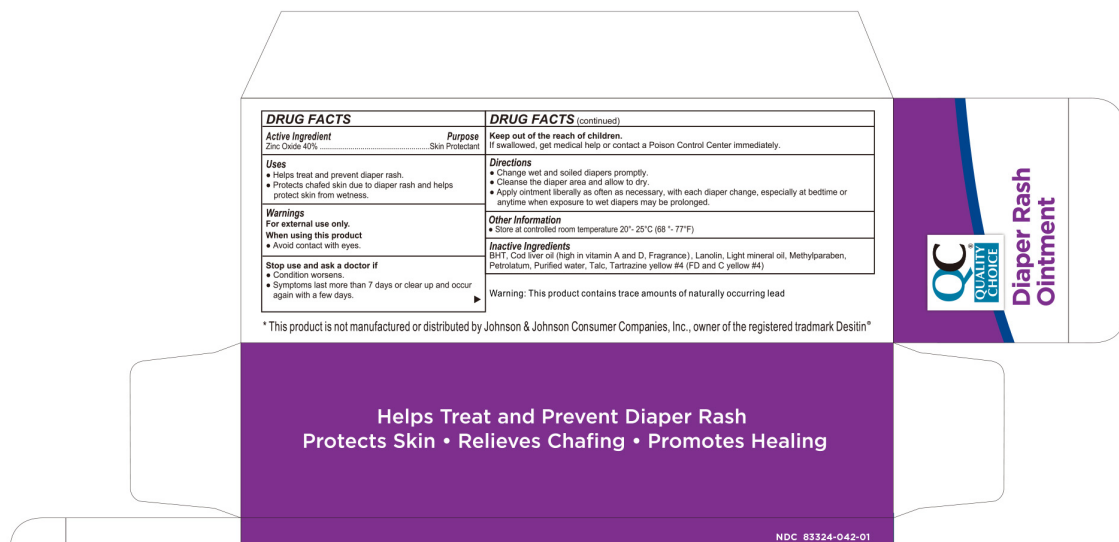
www.qualitychoice.com

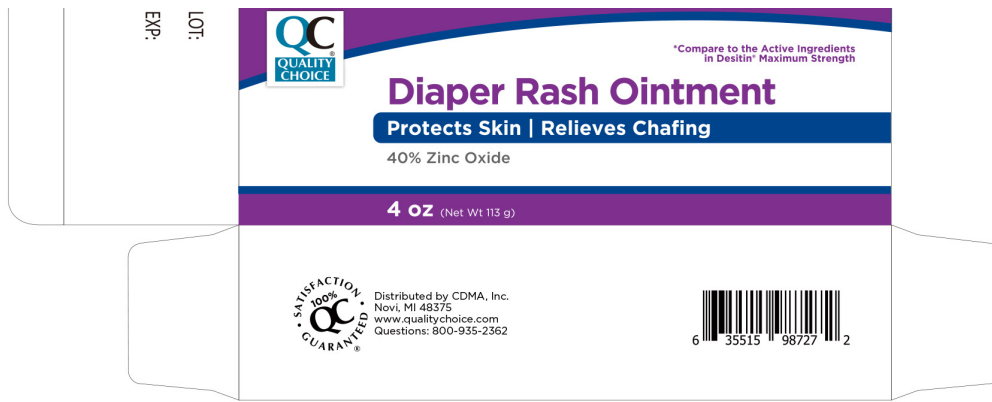
Questions: 800-935-2362

Other Information

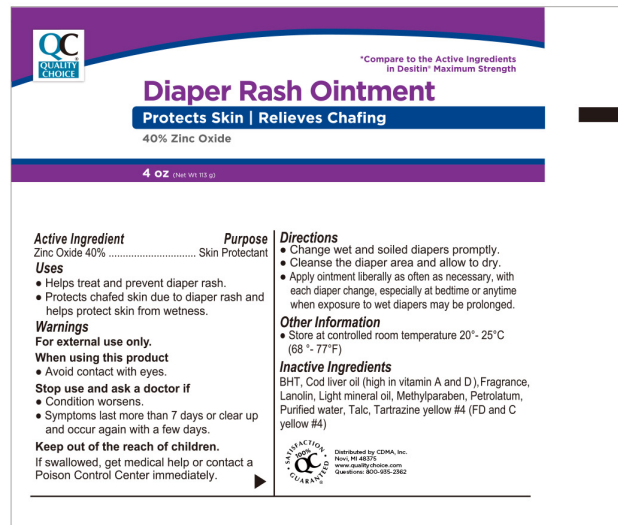
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OUTSIDE BOX





INNER TUBE



QUALITY CHOICE DIAPER RASH

zinc oxide ointment

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:83324-042 |
| Route of Administration | 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: BBL281NWFQ) | |
| 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:83324-042-01 | 113 g in 1 TUBE; Type 0: Not a Combination Product | 05/31/2024 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M016 | 05/31/2024 | |

Labeler - Chain Drug Marketing Association Inc. (011920774)

Registrant - Trifecta Pharmaceuticals USA LLC (079424163)

Revised: 11/2024

Chain Drug Marketing Association Inc.