DE LA CRUZ DIAPER RASH- allantoin, petrolatum, and zinc oxide ointment
DLC Laboratories, 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.

DE LA CRUZ ®DIAPER RASH OINTMENT

Drug Facts

Active ingredients	***Purpose***
Allantoin 1%	Skin protectant
Petrolatum 45%	Skin protectant
Zinc oxide 40%	Skin protectant

Uses

- 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 for more than 7 days or clear up and occur again within a few days

Keep out of the 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 at room temperature

Inactive ingredients

alpha-bisabolol, cholecalciferol, lanolin, lavender oil, phenoxyethanol, retinyl palmitate, tocopheryl acetate, purified water, zea mays (corn) starch

Questions

1-800-858-3889

Manufactured by:
De La Cruz Products
A Division of DLC Laboratories, Inc.
Paramount, CA 90723 USA

PRINCIPAL DISPLAY PANEL - 96 g Tube Label

De La Cruz ®

Baby

Diaper Rash
Ointment

3 in 1
Soothes, Heals, Protects
MAXIMUM
STRENGTH

- Zinc Oxide 40%
- Mild enough for everyday use
- No Artificial Fragrances or Colors,
  No Parabens or Phthalates
- Allergy Tested

NET WT 3.4 OZ (96 g)

DE LA CRUZ DIAPER RASH

allantoin, petrolatum, and zinc oxide ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24286-1566
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)	ALLANTOIN	1 g in 100 g
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	45 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	40 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
LANOLIN (UNII: 7EV65EAW6H)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

STARCH, CORN (UNII: O8232NY3SJ)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
LEVOMENOL (UNII: 24WE03BX2T)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24286-1566-2	1 in 1 CARTON	07/10/2015	11/12/2020
1		48 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:24286-1566-3	96 g in 1 TUBE; Type 0: Not a Combination Product	12/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	07/10/2015	

Labeler - DLC Laboratories, Inc. (093351930)

Establishment

Name	Address	ID/FEI	Business Operations
DLC Laboratories, Inc.		093351930	manufacture(24286-1566) , label(24286-1566)

Revised: 6/2023 DLC Laboratories, Inc.