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

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 Drug	M016	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: 1/2026 DLC Laboratories, Inc.