DESENEX- miconazole nitrate powder
Crown Laboratories

Desenex

Active ingredient

Miconazole nitrate 2%

Purpose

Antifungal

Uses

- Cures most athlete’s foot (tinea pedis)
- Relieves itching, burning, cracking, and discomfort

Warnings

For external use only.

Avoid contact with eyes.

Do not use

on children under 2 years of age unless directed by a doctor.

If irritation occurs or if there is no improvement within 4 weeks, discontinue use and consult a doctor.

Keep out of reach of children.

If swallowed, get medical help or contact a poison control center right away.

Directions

Wash the affected area and dry thoroughly. Apply a thin layer of the product over affected area twice daily (morning and night) or as directed by a doctor. Supervise children in the use of this product. Pay special attention to spaces between the toes; wear well fitting, ventilated shoes, and change shoes and socks at least once daily. Use daily for 4 weeks. If condition persists longer,
consult a doctor. This product is not effective on the scalp or nails.

Other information

- store at 20 º - 25 ºC (68 º - 77 ºF) [see USP Controlled Room Temperature].

Inactive ingredients

1,2-Hexanediol, Aloe Barbadensis Leaf Juice Powder, Beta-Glucan, Caprylyl Glycol, Fragrance, Glycerin, Potassium Sorbate, Sodium Benzoate, Tapioca Starch, Tricalcium Phosphate, Water, Zea Mays (Corn) Starch

Questions?

call **1-833-279-6522**

Principal Display

New & Improved

Desenex®

Antifungal Foot Powder

with 2% Miconazole Nitrate

PRESCRIPTION STRENGTH

Cures Most Athlete's Foot

Triple Action Powder

Relieves Itching, Burning, and Scaling

Attacks and Absorbs Moisture

All Day Odor Control

NET WT. 1.5 oz [43g]

PRODUCT PACKAGED BY WEIGHT NOT VOLUME

Distributed by: Crown Laboratories, Inc **.** Johnson City, TN 37604

P12319.00

DESENEX

miconazole nitrate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0316-0225
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	20 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
STARCH, TAPIOCA (UNII: 24SC3U704I)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
WATER (UNII: 059QF0KO0R)	
STARCH, CORN (UNII: O8232NY3SJ)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
YEAST .BETA.-D-GLUCAN (UNII: 44FQ49X6UN)	
GLYCERIN (UNII: PDC6A3C0OX)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0225-01	43 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/19/2023	
2	NDC:0316-0225-02	85 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/19/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	10/19/2023	

Labeler - Crown Laboratories (079035945)

Revised: 11/2023 Crown Laboratories