

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 associated with athlete's foot

Warnings

For external use only · Avoid contact with eyes.

Do Not Use

- for nail or scalp infections

When using this product

- do not get into the eyes

Stop use and ask a doctor if

- irritation occurs or gets worse
- no improvement within 4 weeks

Keep out of reach of children.

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

Directions

- adults and children 2 years and older
- wash the affected area with soap and water and dry thoroughly before applying
- apply a thin layer over affected area twice a day (morning and night) or as directed by a doctor
- supervise children in the use of this product
- pay special attention to the spaces between the toes. Wear well-fitting, ventilated shoes and change shoes and socks at least once a day.
- use every day for 4 weeks
- children under 2 years of age: ask a doctor

Other information

- Store at 20 ° - 25 °C (68 ° - 77 °F) [see USP Controlled Room Temperature].
- See container bottom for lot number and expiration date

Inactive ingredients

Aldioxa, Chloroxlenol, Croscarmellose Sodium, Fragrance, Imidazolidinyl Urea, Magnesium Stearate, Microcrystalline Cellulose, Sodium Acrylates Crosspolymer-2

Questions or comments?

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

Principal Display

NDC 0316-0225-30

New & Improved

Desenex®

Miconazole Nitrate 2% Antifungal Treatment

PRESCRIPTION STRENGTH

TRIPLE ACTION POWDER

Absorbs Wetness

Repels Moisture

Relieves Itching & Burning

Cures Most Athlete's Foot

NET WT. 3 oz (85 g)

PRODUCT PACKAGED BY WEIGHT NOT VOLUME

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

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P11528.01

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Made in USA

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Drug Facts

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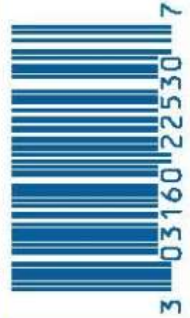
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DESENEX

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: VV4H1CYW1K) (MICONAZOLE - UNII: 7NNO0D7S5M)	MICONAZOLE NITRATE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CHLOROXYLENOL (UNII: 0F32U78V2Q)	
IMIDUREA (UNII: M629807ATL)	
ALDIOXA (UNII: 8T66I31YNK)	
SODIUM ACRYLATES CROSSPOLYMER-2 (UNII: D3HPR4WW6F)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0225-15	43 g in 1 CONTAINER; Type 0: Not a Combination Product	12/01/2018	03/31/2025
2	NDC:0316-0225-30	85 g in 1 CONTAINER; Type 0: Not a Combination Product	12/01/2018	07/24/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	08/03/2009	07/24/2025

Labeler - Crown Laboratories (079035945)

Revised: 12/2023

Crown Laboratories