DESENEX ANTIFUNGAL FOOT CREAM- miconazole nitrate cream
Crown Laboratories, Inc.

Desenex Antifungal Foot Cream

Active ingredient

Miconazole nitrate 2%

Purpose

Antifungal

Uses

- Cures most athlete's foot (tinea pedis)
- Relieves ithcing, burning, cracking, scaling, 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.

Stop use

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-fiting, 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°–25°C (68°–77°F) [see USP Controlled Room Temperature].

Inactive ingredients

Aqua, Benzyl Benzoate, Cetearyl Alcohol, Cetyl Palmitate, Dipropylene Glycol, Fragrance, Paraffinum Liquidum, Polysorbate 60, Potassium Sorbate, Propylene Glycol, Sorbic Acid, Sorbitol, Triethyl Citrate

Questions?

call 1-833-279-6522

Desenex Cream Tube

Desenex

Antifungal Foot Cream

with 2% Miconazole Nitrate

PRESCRIPTION STRENGTH

Cures Most Athlete's Foot

Dual Action Cream

Relieves Itching, Burning, Scaling, and Discomfort

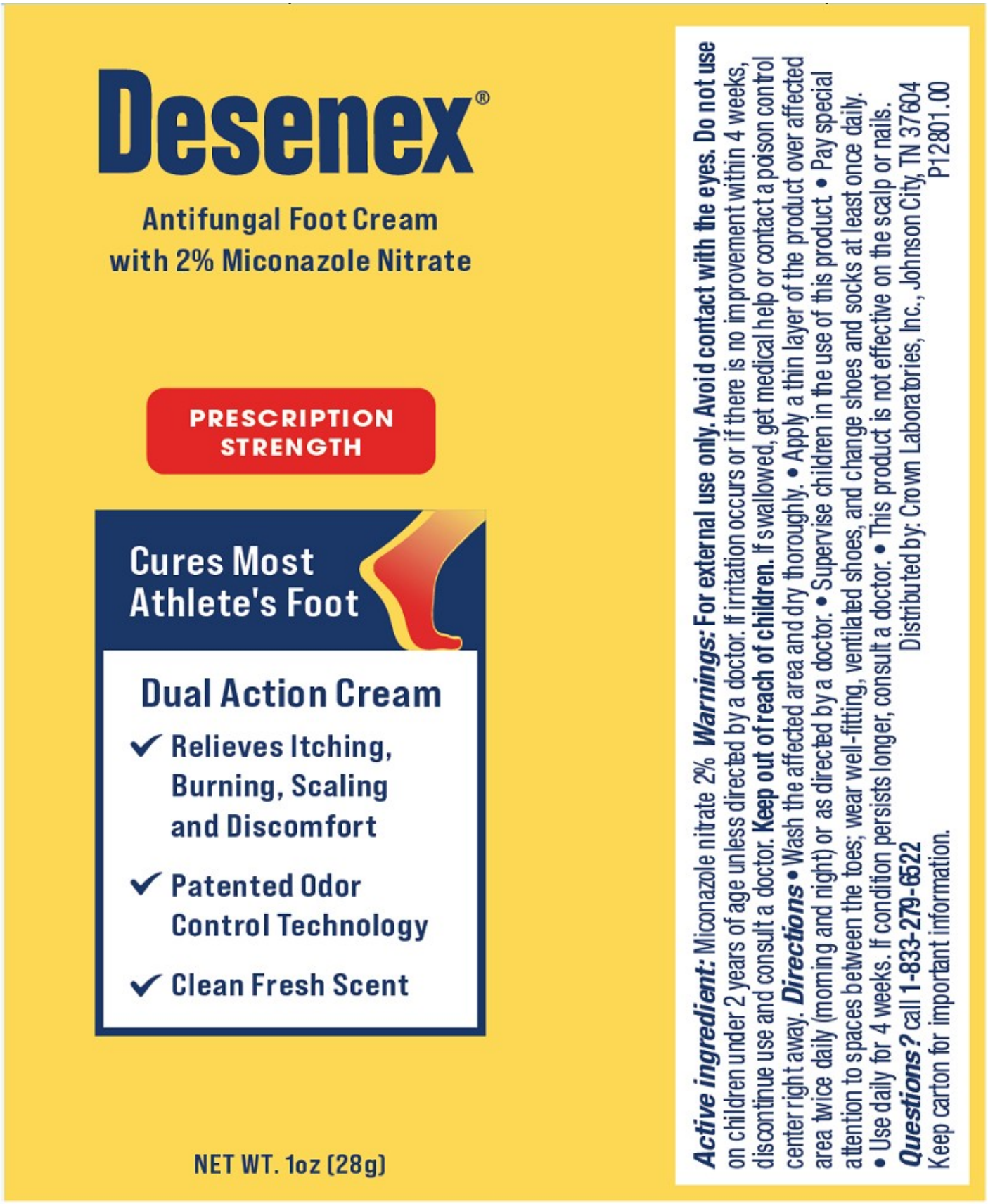
Patented Odor Control Technology

Clean Fresh Scent

NET WT. 1oz (28g)

Distributed by: Crown Laboratories, Inc. Johnson City, TN 36704

P12801.00



Desenex Cream Carton

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Clean Fresh Scent

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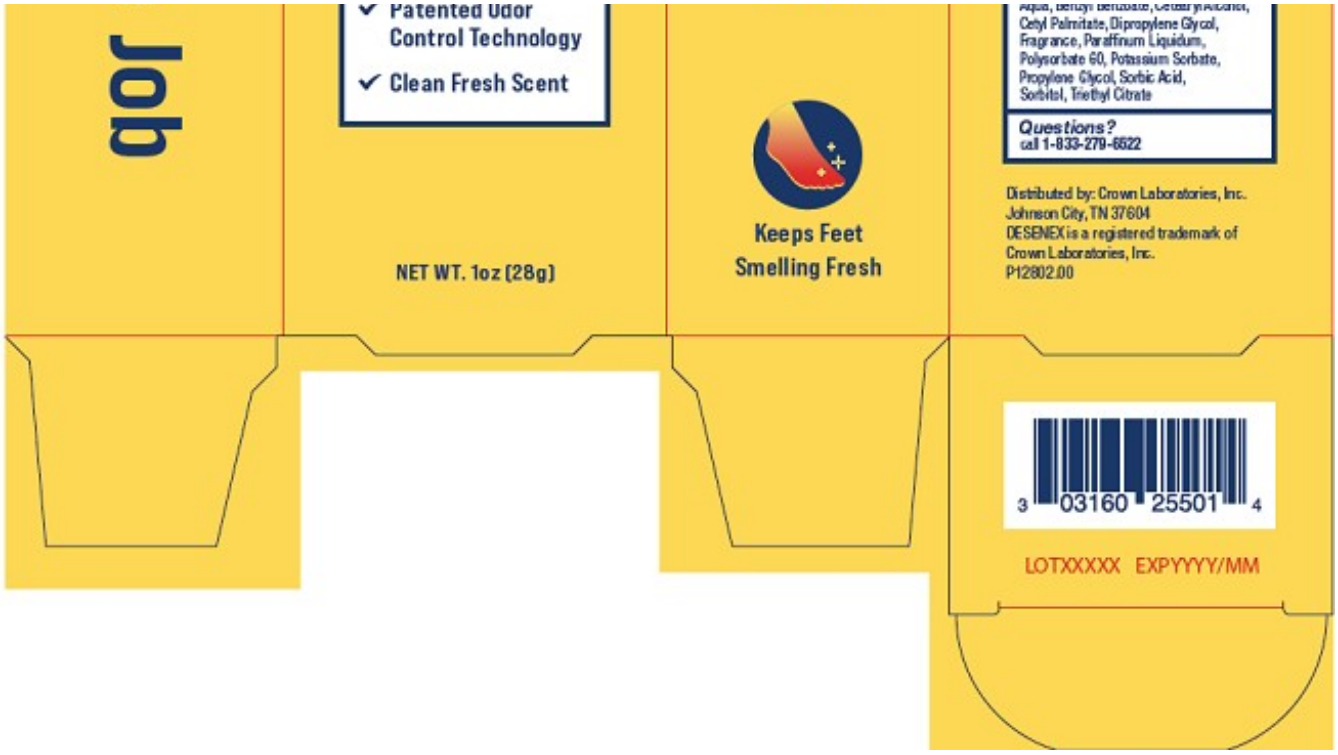
Desenex is a registered trademark of Crown Laboratories, Inc.

P12802.00



DESENEX ANTIFUNGAL FOOT CREAM

miconazole nitrate cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0316-0255
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**
PARAFFINUM LIQUIDUM (UNII: T5L8T28FGP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBIC ACID (UNII: X045WJ989B)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
CETYL PALMITATE (UNII: 5ZA2S6B08X)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
TRIETHYL CITRATE (UNII: 8Z96QXD6UM)	
SORBITOL (UNII: 506T60A25R)	
BENZYL BENZOATE (UNII: N863NB338G)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0255-01	1 in 1 CARTON	03/01/2025	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	03/01/2025	

Labeler - Crown Laboratories, Inc. (079035945)

Establishment

Name	Address	ID/FEI	Business Operations
Crown Laboratories, Inc.		079035945	manufacture(0316-0255)

Revised: 3/2025 Crown Laboratories, Inc.