## MICONAZOLE NITRATE- miconazorb antifungal powder talc free powder Chain Drug Consortium, LLC

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## **Premier Value Miconazorb Antifungal Powder**

### Active ingredient

Miconazole nitrate 2%

## Purpose

**Antifungal** 

#### Uses

for the cure of most athlete's foot, jock itch and ringworm

#### Warnings

For external use only.

#### Do not use

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

## When using this product

avoid contact with the eyes

## Stop and ask a doctor if

irritation occurs or there is no improvement within 4 weeks for athlete's foot and ringworm, or 2 weeks for jock itch.

## Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away. Do not use on children under 2 years of age unless directed by a doctor.

#### **Directions**

- clean the affected area and dry thoroughly
- apply a thin layer of product over affected area twice daily (morning and night) or as directed by a doctor
- supervise children in the use of this product
- for athlete's foot: pay special attention to spaces between the toes; wear well-fitting, ventilated shoes, and change shoes and socks at least once daily
- for athlete's foot and ringworm, use daily for 4 weeks; for jock itch, use daily for 2 weeks

- if conditions persist longer, consult a doctor
- this product is not effective on the scalp or nails

#### Other information

- store between 59º 86ºF
- lightly shake bottle to loosen settled powder

## Inactive ingredients

allantoin, chloroxylenol, fragrance, imidazolidinyl urea, microcrystalline cellulose, tricalicum phosphate, zea mays (corn) starch

### Questions?

call 1-866-964-0939

## **Principal Display Panel**

**Premier Value** 

Miconazorb AF

#### ANTIFUNGAL POWDER

Miconazole Nitrate 2%

Cures most athlete's foot, jock itch & ringworm

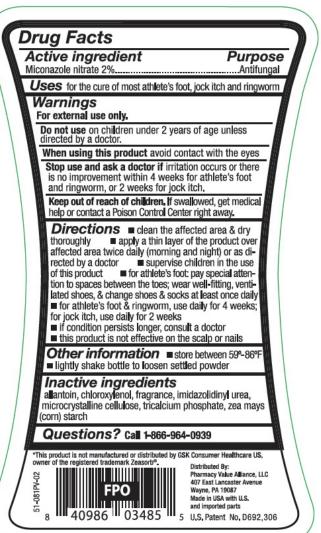
Relieves chafing, itching, burning & scaling

Absorbs moisture

Talc-free

**NET WT 2.5 OZ (71 g)** 





#### MICONAZOLE NITRATE

miconazorb antifungal powder talc free powder

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:68016-673 Route of Administration TOPICAL

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M) MICONAZOLE NITRATE | 1.42 g in 71 g

Inactive Ingredients			
Ingredient Name	Strength		
CHLOROXYLENOL (UNII: 0F32U78V2Q)			

ALLANTOIN (UNII: 344S277G0Z)	
IMIDUREA (UNII: M629807ATL)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
ZEA MAYS SUBSP. MAYS WHOLE (UNII: 1G5HNE09V8)	

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:68016- 673-25	71 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/22/2017	01/30/2026		

<b>Marketing In</b>	larketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M005	10/22/2017	01/30/2026		

## Labeler - Chain Drug Consortium, LLC (101668460)

Revised: 12/2024 Chain Drug Consortium, LLC