MICONAZOLE NITRATE- miconazole nitrate powder
AvKARE

Miconazole Nitrate 2%

Drug Facts

Active ingredient

Miconazole Nitrate 2.0%

Purpose

Antifungal

Uses

- for the treatment of most athlete's foot (tinea pedis), jock itch (tinea cruris), ringworm (tinea corporis)
- for the treatment of most superficial skin infections caused by yeast (candida albicans)
- relieves most itching, scaling, cracking, burning, redness, soreness, irritation, discomfort and chafing associated with jock itch

Warnings

- **for external use only**

Do not use

- on children under 2 years of age unless by a doctor
- avoid contact with the eyes
- for athlete's foot and ringworm - if irritation occurs, or if there is no improvement within 4 weeks, discontinue use and consult a doctor
- for jock itch - if irritation occurs, or if there is no improvement within 2 weeks, dicontinue 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

- clean the affected area and dry thoroughly
- apply a thin layer of powder 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 toes; wear well - fitting, ventilated shoes, and change shoes and socks at least once daily
- for athletes foot and ringworm, use daily for 4 weeks

- for jock itch, use daily for 2 weeks
- if condition persists longer, consult a doctor
- this product is not effective on the scalp or nails

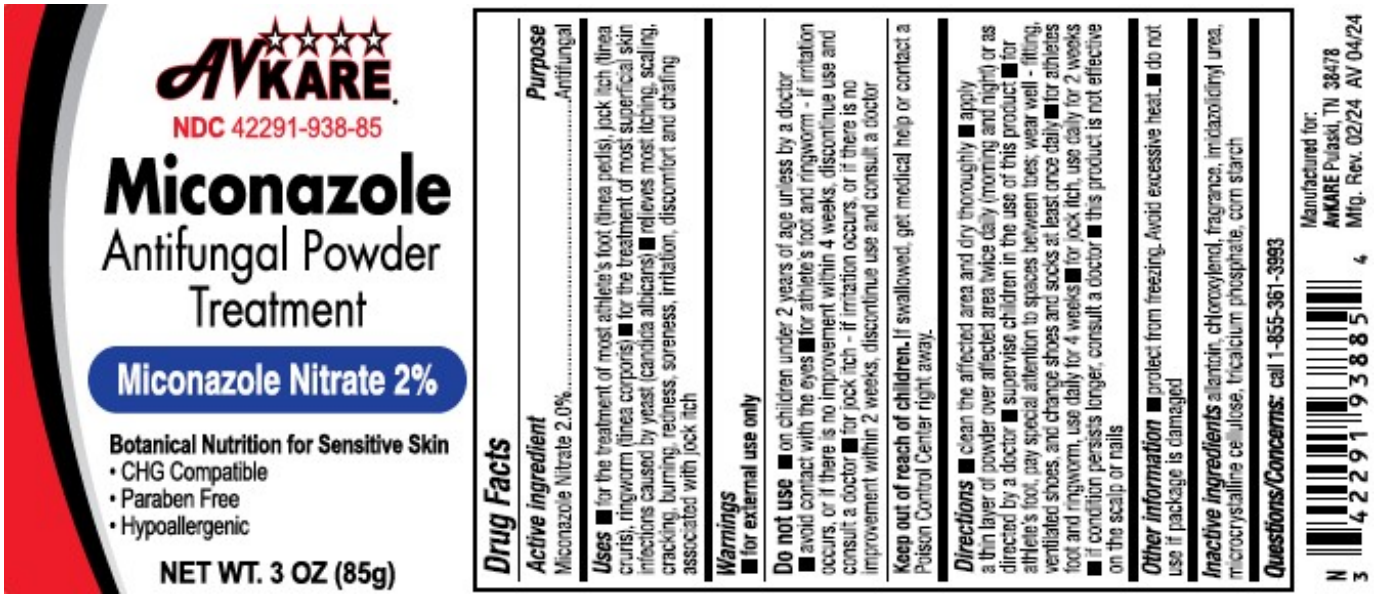
Other information

- protect from freezing. Avoid excessive heat.
- do not use if package is damaged

Inactive ingredients

allantoin, chloroxylenol, fragrance, imidazolidinyl urea, microcrystalline cellulose, tricalcium phosphate, corn starch

PRINCIPAL DISPLAY PANEL - 85 g Bottle Label



MICONAZOLE NITRATE

miconazole nitrate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42291-938(NDC:69367-399)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	2 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**

ALLANTOIN (UNII: 344S277G0Z)	
CHLOROXYLENOL (UNII: 0F32U78V2Q)	
IMIDUREA (UNII: M629807ATL)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white (White to off-white)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42291-938-85	85 g in 1 BOTTLE; Type 0: Not a Combination Product	06/05/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	06/05/2024	

Labeler - AvKARE (796560394)

Revised: 6/2024 AvKARE