

**MICONAZORB AF- miconazole nitrate powder**  
**AmerisourceBergen Drug Corporation**

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**Good Neighbor Pharmacy Miconazorb AF**

***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 under 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.

***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 59° - 86°F
- lightly shake bottle to loosen settled powder

***Inactive ingredients***

Zea Mays (Corn) Starch, Tricalcium Phosphate, Microcrystalline Cellulose, Sodium Bicarbonate, Allantoin, Chloroxylonol, Fragrance

***Questions?***

call 1-866-964-0939

**Principal Display Panel**

Good Neighbor Pharmacy

- Talc-Free

Miconazorb AF

Miconazole Nitrate 2%/Antifungal

- Cures most athlete's foot, jock itch & ringworm
- Relieves itching, burning, scaling & chafing associated with jock itch
- Absorbs moisture
- Comforts & refreshes

NET WT. 2.5 OZ. (71 g)



## MICONAZORB AF

miconazole nitrate powder

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46122-789
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
ALLANTOIN (UNII: 344S277G0Z)	
CHLOROXYLENOL (UNII: 0F32U78V2Q)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STARCH, CORN (UNII: O8232NY3SJ)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-789-29	71 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/05/2024	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	04/05/2024	

**Labeler** - AmerisourceBergen Drug Corporation (007914906)

Revised: 10/2025

AmerisourceBergen Drug Corporation