MICONAZOLE NITRATE- anti-fungal powder miconazole nitrate talc free powder
AmerisourceBergen Drug Corporation

GNP Miconazorb Antifungal Powder Talc-Free

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

GOOD NEIGHBOR PHARMACY

Miconazorb AF

Miconazole Nitrate 2%

ANTIFUNGAL POWDER

- Cures most athlete's foot, jock itch and ringworm
- Relieves chafing, itching, burning and scaling
- Absorbs moisture
- Talc-free

NET WT 2.5 OZ (71 g)

MICONAZOLE NITRATE

anti-fungal powder miconazole nitrate talc free powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:46122-444
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-444-27	71 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/08/2017	05/31/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	08/08/2017	05/31/2026

Labeler - AmerisourceBergen Drug Corporation (007914906)

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