

MICONAZOLE NITRATE- anti-fungal powder miconazole nitrate talc free powder
Hudson Health LLC

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

Inactive ingredients

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

Questions?

Call 1-877-237-0194

Principal Display Panel

comfortzone

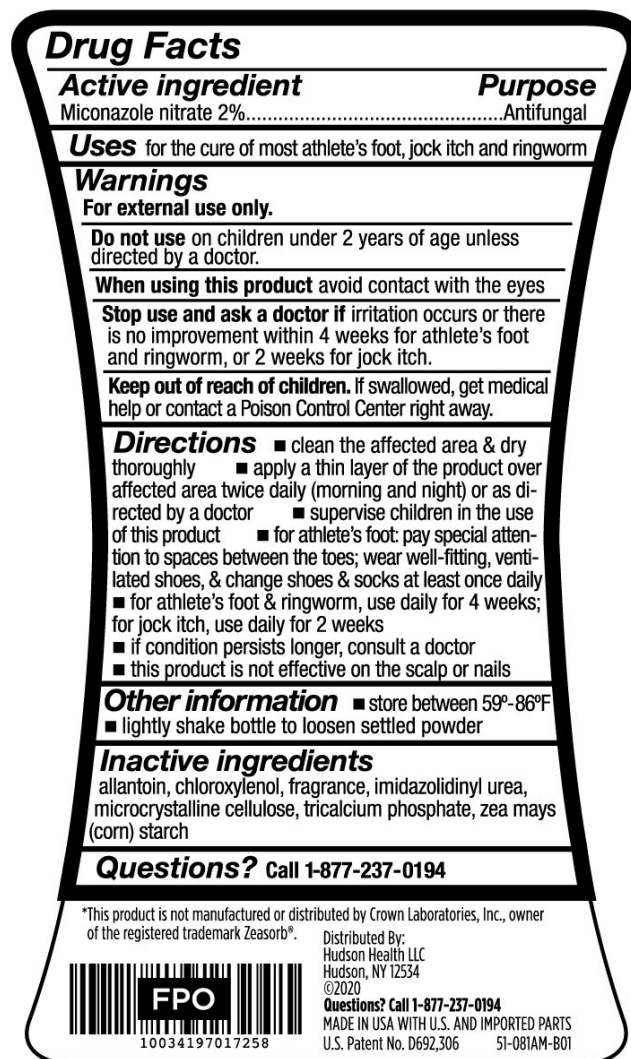
miconazorbAF

antifungal powder

micronazole nitrate

- cures most athlete's foot, jock itch & ringworm
- relieves chafing, itching, burning & scaling
- absorbs moisture
- talc-free

NET WT 2.5 OZ (71g)



MICONAZOLE NITRATE

anti-fungal powder miconazole nitrate talc free powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72446-007
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
ALLANTOIN (UNII: 344S277G0Z)	
CHLOROXYLENOL (UNII: 0F32U78V2Q)	

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:72446-007-71	71 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	09/01/2018	

Labeler - Hudson Health LLC (081276171)

Revised: 1/2024

Hudson Health LLC