MEDICATED ATHLETES FOOT POWDER- miconazole nitrate powder
Premier Brands of America Inc.

Arm & Hammer Medicated Athlete's Foot 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 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 the 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, & change shoes & socks at least once daily
- for athlete's foot & 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

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

Questions?

call 1-866-964-0939

Principal Display Panel

ARM & HAMMER

MEDICATED

ATHLETE'S

FOOT

POWDER

MICONAZOLE NITRATE 2% /

ANTIFUNGAL TREATMENT

CURES MOST

ATHLETE'S FOOT,

JOCK ITCH &

RINGWORM

Relieves Itching, Burning,

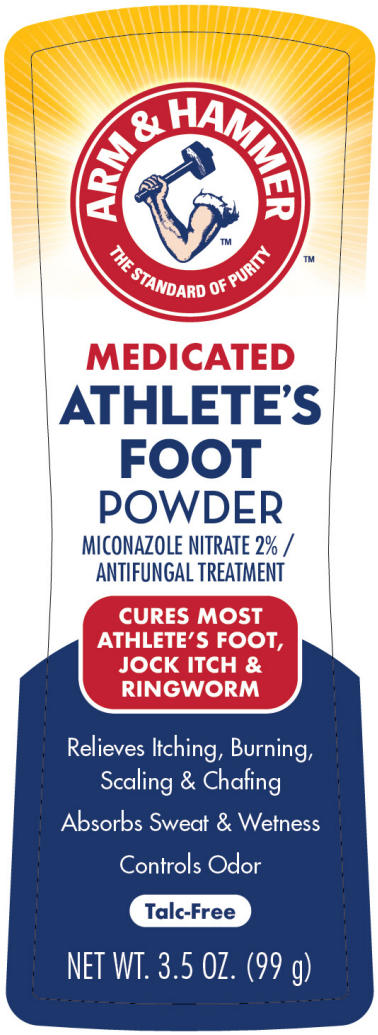
Scaling & Chafing

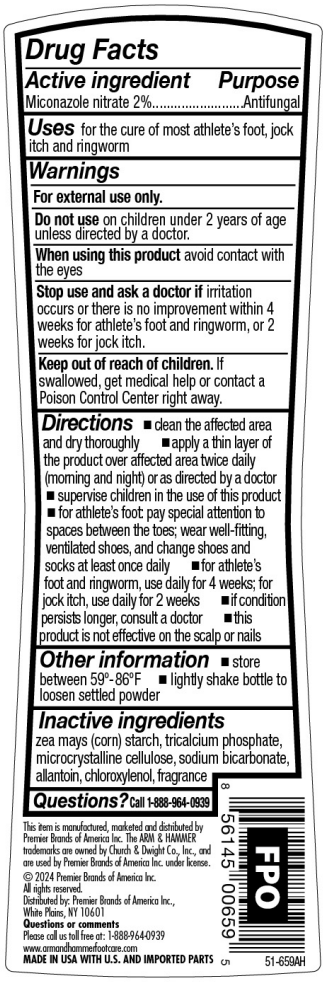
Absorbs Sweat & Welness

Controls Odor

Talc-free

NET WT. 3.5 OZ. (99 g)





MEDICATED ATHLETES FOOT POWDER

miconazole nitrate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:56104-906
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE -	MICONAZOLE NITRATE	20 mg in 1 g

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:56104-906-01	99 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/26/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	06/24/2025	

Labeler - Premier Brands of America Inc. (063849780)

Revised: 6/2025 Premier Brands of America Inc.