

LOTRIMIN ANTIFUNGAL- miconazole nitrate powder
Bayer HealthCare LLC.

Lotrimin®

Antifungal

Drug Facts

Active ingredient

Miconazole nitrate 2%

Purpose

Antifungal

Uses

- Cures most athlete's foot (tinea pedis), jock itch (tinea cruris) and ringworm (tinea corporis)
- relieves itching, cracking, burning and scaling

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 use and ask a doctor if

- irritation occurs
- if 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

- wash affected area and dry thoroughly
- sprinkle a thin layer over affected area twice daily (morning and night)
- 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 condition persists longer, ask a doctor
- this product is not effective on the scalp or nails

Other information

store between 20° to 25°C (68° to 77°F)

Inactive ingredients

benzethonium chloride, corn starch, kaolin, sodium bicarbonate, starch/acrylates/acrylamide copolymer, zinc oxide

PRINCIPAL DISPLAY PANEL - 90g Can Label

NDC 11523-0919-1

LOTRIMIN[®] AF

ANTIFUNGAL

miconazole nitrate

Cures Most

Athlete's Foot

POWDER

- **Absorbs Moisture**
- **Relieves Itching,**

Burning & Scaling

NET WT 90g (3 OZ)

37871-00

LOTTRIMIN[®]
MICONAZOLE NITRATE **ANTIFUNGOAL**

MEDICATED FOOT POWDER

CLINICALLY PROVEN TO CURE MOST
ATHLETE'S FOOT

RELIEVES

- ITCHING
- BURNING
- SCALING

Absorbs moisture



C00012114

NET WT 90g (3 OZ)



LOTRIMIN ANTIFUNGAL

miconazole nitrate powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11523-0919
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
STARCH, CORN (UNII: O8232NY3SJ)	
KAOLIN (UNII: 24H4NWX5CO)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
ZINC OXIDE (UNII: SOI2LOH54Z)	

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:11523-0919-1	90 g in 1 CAN; Type 0: Not a Combination Product	09/01/1993	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	09/01/1993	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 1/2026

Bayer HealthCare LLC.