SILKA ATHLETES FOOT- miconazole nitrate aerosol, spray
Genomma Lab USA

Silka® Athlete's Foot Spray

Drug Facts

Active ingredient

Miconazole nitrate 2%

Purpose

Antifungal

Uses

- Clinically proven to be effective in the treatment of most athlete's foot (tinea pedis)
- Provides effective relief of itching, cracking, burning, scaling, and discomfort.

Warnings

For external use only.

Flammable. Do not use while smoking or near heat or flame.

Do not use on children under 2 years of age unless directed by a doctor.

When using this product

- Avoid contact with eyes
- in case of contact with eyes, flush thoroughly with water
- Avoid applying on broken skin
- do not inhale or ingest
- contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120°F

Stop use and ask a doctor if

- irritation occurs
- there is no improvement within 4 weeks (for athlete's foot)

Keep out of the reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- wash affected area and dry thoroughly
- shake can well and spray 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
- this product is not effective on the scalp or nails

Other information
- store between 20° to 25° (68° to 77°F)

Inactive ingredients

Alcohol Denat., polysorbate 20, tetrafluoropropene

Questions?

1-877-994-3666 Monday to Friday, 8am to 6pm Central time.

Distributed by: Genomma Lab, Houston, TX 77098

PRINCIPAL DISPLAY PANEL - 130 g Canister Label

SILKA®
ATHLETE'S FOOT
POWDER SPRAY

Miconazole nitrate 2% - antifungal

CLINICALLY PROVEN TO CURE MOST
ATHLETE'S FOOT*

HELPS RELIEVE
- BAD ODOR
- ITCHING
- SWEATING
- BURNING

Net Wt 4.6oz (130g)

SILKA ATHLETES FOOT

miconazole nitrate aerosol, spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50066-084
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**
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
ALCOHOL (UNII: 3K9958V90M)	
1-CHLORO-3,3,3-TRIFLUOROPROPENE (UNII: 4B96DT2BYB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50066-084-46	130 g in 1 CANISTER; Type 0: Not a Combination Product	02/03/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M005	02/03/2025	

Labeler - Genomma Lab USA (832323534)

Revised: 2/2025

Genomma Lab USA