

**ANTIFUNGAL POWDER- miconazole nitrate powder  
LEADER/ Cardinal Health 110, Inc.**

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**Leader Antifungal Powder**

***Active ingredient***

Miconazole nitrate 2%

***Purpose***

Antifungal

***Uses***

for the cure of most athlete's foot, ringworm and chafing associated with jock itch

***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 59° - 86°F
- 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**

LEADER

Moisture Absorbing

Antifungal

Powder

Miconazole Nitrate 2%/

Antifungal

Cures Most Athlete's Foot

Relieves Itching, Burning

& Scaling

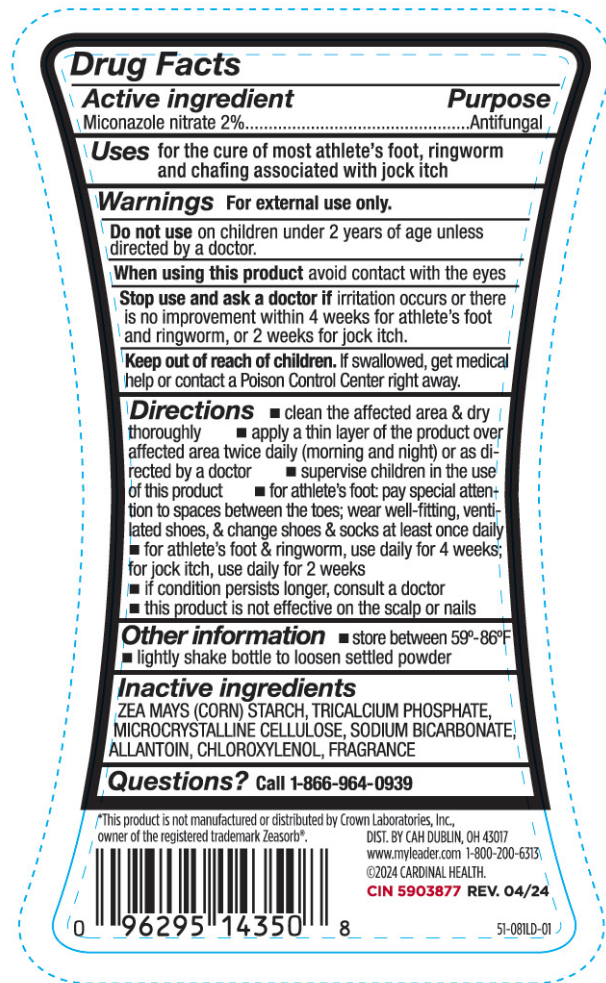
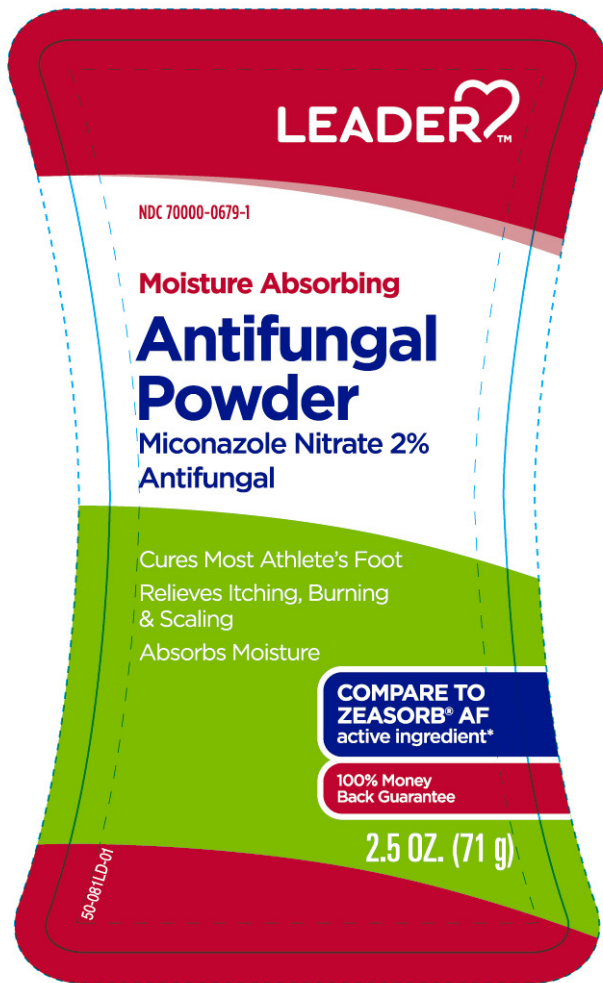
Absorbs Moisture

COMPARE TO

ZEASORB AF

active ingredient

NET WT 2.5 OZ (71 g)



## ANTIFUNGAL POWDER

miconazole nitrate powder

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70000-0679
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>MICONAZOLE NITRATE</b> (UNII: VV4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	20 mg in 1 g

### Inactive Ingredients

Ingredient Name	Strength
<b>ALLANTOIN</b> (UNII: 344S277G0Z)	
<b>CHLOROXYLENOL</b> (UNII: 0F32U78V2Q)	

<b>SODIUM BICARBONATE</b> (UNII: 8MDF5V39QO)	
<b>TRICALCIUM PHOSPHATE</b> (UNII: K4C08XP666)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0679-1	71 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/16/2024	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	08/16/2024	

**Labeler** - LEADER/ Cardinal Health 110, Inc. (063997360)

Revised: 11/2024

LEADER/ Cardinal Health 110, Inc.