

MICONAZOLE NITRATE 2%- miconazole nitrate cream
Akron Pharma Inc.

MEDPURA MICONAZOLE Nitrate 2%
ANTIFUNGAL CREAM

Soothes itching, scaling, cracking and burning
Cures most athlete's foot, jock itch and ringworm

Drug Facts

Active ingredient

Miconazole Nitrate, USP 2%

Purpose

Anti-Fungal

Uses

- proven clinically effective in the treatment of most athlete's foot, jock itch, and ringworm
- for effective relief of itching, scaling, cracking, burning, and discomfort

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 and ringworm, use daily for 4 weeks; for jock itch, use daily for 2 weeks
- if condition lasts longer, contact a doctor + this product is not effective on the scalp or nails

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
- there is no improvement within 4 weeks (for athlete's foot and ringworm) or within 2 weeks (for jock itch)

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Other Information

- do not use if foil seal on tube opening is broken or missing
- to open tube: unscrew cap, lift tab, and pull to remove foil seal prior to use
- store at room temperature 20°-25°C (68°-77°F)
- before using any medication, read all label directions. Keep carton, it contains important information.

Inactive ingredients Cetostearyl Alcohol, EthylParaben, Glycerin, Glyceryl Monostearate, Mineral Oil, Petrolatum, Polyoxyl lauryl ether, Purified water, Stearic acid

Call toll-free 1(877)225-6999

Manufactured for:

Akron Pharma, Inc.

Fairfield, NJ 07004

**Miconazole Nitrate Cream USP, 2%
Antifungal Cream**

Rev.:12/2024

Akron Pharma
Manufactured for:
Akron Pharma, Inc.
Fairfield, NJ 07004

N 3 71399 00711 7

NDC 71399-0071-1

*Compare to the active ingredient in Micatin® Antifungal Cream

**Miconazole Nitrate Cream USP, 2%
Antifungal Cream**

Soothes itching, scaling, cracking and burning
Cures most athlete's foot, jock itch and ringworm

Net wt. 0.5oz(14.2g)

Drug Facts	
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Miconazole Nitrate Cream USP, 2% Antifungal Cream

Akron Pharma
Manufactured for:
Akron Pharma, Inc.
Fairfield, NJ 07004

Rev.:12/2024




MEDPURA

NDC 71399-0071-3

*Compare to the active ingredient in
Micatin® Antifungal Cream

Miconazole Nitrate Cream USP, 2% Antifungal Cream

Soothes itching, scaling, cracking and burning
Cures most athlete's foot, jock itch and ringworm

Net wt. 1oz(28.4g)


MEDPURA

Drug Facts

Active ingredient	Purpose
Miconazole nitrate USP, 2%	Antifungal

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Questions or comments? Call toll-free 1(877)225-6999

MEDPURA


Miconazole Nitrate Cream USP, 2%

Antifungal Cream



Manufactured for:
Akron Pharma, Inc.
Fairfield, NJ 07004

Rev.:12/2024



N 3 71399 00714 8



NDC 71399-0071-4

*Compare to the active ingredient in Micatin® Antifungal Cream

Miconazole Nitrate Cream USP, 2%

Antifungal Cream

Soothes itching, scaling, cracking and burning
Cures most athlete's foot, jock itch and ringworm

Net wt. 1.5oz(42.5g)



Drug Facts

<i>Active ingredient</i> Miconazole nitrate USP, 2%	<i>Purpose</i> Antifungal
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MICONAZOLE NITRATE 2%

miconazole nitrate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71399-0071
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII: 7NNO0D7S5M)	MICONAZOLE NITRATE	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
ETHYLPARABEN (UNII: 14255EXE39)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	

MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
LAURETH-23 (UNII: N72LMW566G)	
WATER (UNII: 059QF0KO0R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-0071-1	1 in 1 BOX	11/19/2024	
1		14.2 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:71399-0071-3	1 in 1 BOX	11/19/2024	
2		28.4 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:71399-0071-4	1 in 1 BOX	11/19/2024	
3		42.5 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	11/19/2024	

Labeler - Akron Pharma Inc. (067878881)

Revised: 1/2025

Akron Pharma Inc.