

JOCK ITCH POWDER- miconazole nitrate powder
Premier Brands of America Inc

Premier Solutions Jock Itch 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, 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 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, MICOCRYSTALLINE CELLULOSE, SODIUM BICARBONATE, ALLANTOIN, CHLOROXYLENOL, FRAGRANCE

Questions?

call 1-866-964-0939

Principal Display Panel

premier

solutions

TALC-FREE

JOCK ITCH

POWDER

MICONAZOLE NITRATE 2% /

ANTIFUNGAL

CURES MOST JOCK ITCH

Relieves itching, burning,

scaling & chafing

Absorbs moisture

NET WT. 2.5 OZ. (71 g)



Drug Facts

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FPO

DISTRIBUTED BY Premier Brands of America Inc. White Plains, NY 10601 QUESTIONS? CALL 1-866-964-0939 MADE IN USA FROM GLOBALLY SOURCED MATERIALS

JOCK ITCH POWDER

miconazole nitrate powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56104-901
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
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:56104-901-71	71 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	06/01/2023	

Labeler - Premier Brands of America Inc (063849780)

Revised: 5/2024

Premier Brands of America Inc