# JOCK ITCH POWDER- miconazole nitrate powder Walgreens Company

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## Walgreens 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 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 59º 86ºF
- lightly shake bottle to loosen settled powder

# Inactive ingredients

allantoin, chloroxylenol, fragrance, micocrystalline cellulose, sodium bicarbonate, tricalcium phosphate, zea mays (corn) starch

#### Questions?

call 1-866-964-0939

# **Principal Display Panel**

Walgreens

Jock

Itch

Powder

MICONAZOLE NITRATE 2% /

ANTIFUNGAL POWDER

Cures most jock itch

Relieves itching, burning,

scaling & chafing

Absorbs moisture

Talc-free

NET WT 2.5 OZ. (71 g)



## Drug Facts Active ingredient **Purpose** 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 use 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 & dry thoroughly apply a thin layer of the product ove affected area twice daily (morning and night) or as directed by a doctor supervise children in the use of this product from a thlete'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 condition persists longer, consult a doctor ■ this product is not effective on the scalp or nails Other information ■ store between 59°-86°l ■ lightly shake bottle to loosen settled powder Inactive ingredients ALLANTOIN, CHLOROXYLENOL, FRAGRANCE, MICROCRYSTALLINE CELLULOSE, SODIUM BICARBONATE, TRICALCIUM PHOSPHATE, ZEA MAYS (CORN) STARCH. Questions? Call 1-866-964-0939

# JOCK ITCH POWDER

miconazole nitrate powder

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-9223

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name

MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)

Basis of Strength

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: 08232NY3SJ)

Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:0363- 9223-01	71 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/09/2024				
Marketing Information							

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M005	05/09/2024			

# Labeler - Walgreens Company (008965063)

Revised: 5/2024 Walgreens Company