

MICONAZOLE NITRATE- antifungal powder
TWIN MED, LLC

Anti-Fungal Powder Talc

Active Ingredients

Miconazole nitrate 2%

Purpose

Antifungal

Uses

For the treatment of most athlete's foot (tinea pedis), jock itch (tinea cruris), ringworm (tinea corporis) ■Relieves itching, scaling, cracking, burning, redness, soreness, irritation, discomfort and chafing associated with jock itch or itching, burning feet

Warnings

For external use only.

Do not use

: ■On children under 2 years of age unless directed by a doctor ■For diaper rash

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

■Irritation occurs

■There is no improvement within 2 weeks when used for the treatment of jock itch

■There is no improvement within 4 weeks when used for athlete's foot or ringworm

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 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 condition persists longer, consult a doctor
 - this product is not effective on the scalp or nails

Other information

- Store at room temperature

Inactive ingredients

Colloidal Anhydrous Silica, Talcum Powder

Pincipal Display Panel

PROCURE™
HEALTHCARE PRODUCTS

ANTIFUNGAL POWDER

MICONAZOLE NITRATE 2%

*Helps treat foot odor,
athlete's foot
and ringworm*

NET WT.: 3 OZ. (85G)



Reorder No. PCFP03

Drug Facts

Active ingredient	Purpose
Miconazole Nitrate (2% w/w).....	Antifungal

Uses ■ For the treatment of most athlete's foot (tinea pedis), jock itch (tinea cruris), ringworm (tinea corporis) ■ Relieves itching, scaling, cracking, burning, redness, soreness, irritation, discomfort and chafing associated with jock itch or itching, burning feet

Warnings: For external use only.

Do not use: ■ On children under 2 years of age unless directed by a doctor ■ For diaper rash

Ask a doctor or a pharmacist before use if you are taking: ■ Anti coagulants ■ Warfarin

When using this product avoid contact with eyes

Stop use and ask a doctor if ■ Irritation occurs ■ There is no improvement within 2 weeks when used for the treatment of jock itch ■ There is no improvement within 4 weeks when used for athlete's foot or ringworm

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 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 condition persists longer, consult a doctor ■ This product is not effective on the scalp or nails

Other information: Store at room temperature

Inactive ingredients:

Colloidal Anhydrous Silica, Talcum Powder

LOT



Distributed by
Twin Med LLC
Santa Fe Springs, CA 90670
www.procureproducts.com

Made in India
HR/DRUGS 1105-OSP(H)



MICONAZOLE NITRATE

antifungal powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55681-165
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TALC (UNII: 7SEV7J4R1U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55681-165-03	85 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/19/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	08/19/2022	

Labeler - TWIN MED, LLC (009579330)

Revised: 6/2025

TWN MED, LLC