MICONAZOLE NITRATE- antifungal powder
TWIN MED, LLC

Anti-Fungal Powder Talc

Active Ingredients

Miconazole nitrate 2%

Purpose

Antifungal

Uses

fFor 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 Anhyhdrous 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)

PROCURE PRODUCTS SYMBOL OF EXCELLENCE

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 Anhyhdrous 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)

8 16516 01563 2

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: 1/2024 TWIN MED, LLC