QUINSANA ATHLETES FOOT ANTI FUNGAL- tolnaftate powder
Profoot, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Quinsana Athlete's Foot Anti-Fungal Powder

Drug Facts

Active Ingredient

Tolnaftate, 1%

Purpose

Antifungal

Uses

- Cures most athlete's foot (tinea pedis), jock itch (tinea cruris), and ringworm (tinea corporis)
- Effectively relieves itching, cracking, and burning

Warnings

For external use only

Ask a doctor before use

- on children under 2 years of age

When using this product

- avoid contact with eyes

Stop use and ask a doctor if

- irritation occurs
- 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

- Wash affected area and dry thoroughly.
- Apply a thin layer of the product over time indicated.
- Supervise children in the use of this product. **For athlete's foot and ringworm:** Use twice daily (morning and night) for 4 weeks or as directed by physician. Pay special attention to spaces between the toes. Wear well-fitting, ventilated shoes, and change shoes and socks at least once daily. **For jock itch:** Use daily for 2 weeks.
- Clean the affected area and dry thoroughly.
- This product is not effective on the scalp or nails.

Inactive ingredients

Talc, Cornstarch.

Package Labeling:

QUINSANA ATHLETES FOOT ANTI FUNGAL

tolnaftate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:29784-141
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	10 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
TALC (UNII: 7SEV7J4R1U)	
STARCH, CORN (UNII: O8232NY3SJ)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29784-141-85	85 g in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333C	05/01/2019	

Labeler - Profoot, Inc. (107570900)

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