

**TOLNAFATE- tolnaftate powder spray aerosol, spray**  
**Chain Drug Marketing Association**

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**Quality Choice Tolnaftate Powder Spray Talc-Free**

***Active ingredient***

Tolnaftate 1%

***Purpose***

Antifungal

***Uses***

- cures most athlete's foot (tinea pedis), ringworm (tinea corporis) and jock itch (tinea cruris)
- helps prevent most athlete's foot with daily use
- relieves itching, burning, cracking, and chafing associated with jock itch

**For external use only.**

**Extremely Flammable:**

Keep away from fire, sparks, and heated surfaces.

Contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120°F.

**Do not use**

on children under 2 years of age unless directed by a doctor.

**When using this product**

- avoid spraying in eyes
- use only as directed
- intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal.

**Stop use and ask a doctor if**

- there is no improvement within 4 weeks for athlete's foot and ringworm; 2 weeks for jock itch
- irritation occurs

**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
- shake can well and spray a thin layer of the product over affected area twice daily (morning and night) or as directed by a doctor
- to prevent athlete's foot, shake can well and spray a thin layer of the product to the feet once or twice daily ( morning and/or night)
- 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, consult a doctor
- this product is not effective on the scalp or nails

**Other information**

- store between 20° and 30°C (68° and 86°F)
- if nozzle clogs, clean with a pin

**Inactive ingredients**

Isobutane, SD Alcohol 40-B, Kaolin, Zea Mays (Corn) Starch, BHT, PPG-12-Buteth-16

**Questions?**

call 1-866-964-0939

**Principal Display Panel****QC Quality Choice****Athlete's****Foot****Powder****Spray****Tolnaftate 1% / Antifungal**

Cures Most Athlete's Foot

& Prevents Recurrences

Comforts & Refreshes

Talc-Free

Relieves:

Itching

Burning

Cracking

**NET WT.**

**4.6 OZ. (130 g)**

NDC 83324-234-46



Compare to the  
Active Ingredient In  
Tinactin®\*

# Athlete's Foot Powder Spray

**Tolnaftate 1% /Antifungal**

Cures Most Athlete's Foot  
& Prevents Recurrences

Comforts & Refreshes

Talc-Free

**Relieves:**  
Itching  
Burning  
Cracking



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4.6 OZ. (130 g)**

\*This product is not manufactured or distributed by Bayer HealthCare LLC, owner of the registered trademark Tinactin®.

<b>Drug Facts</b>	
<b>Active ingredient</b>	<b>Purpose</b>
Tolnaftate 1%.....	.....Antifungal
<b>Uses</b> ■ cures most athlete's foot (tinea pedis), ringworm (tinea corporis) and jock itch (tinea cruris) ■ helps prevent most athlete's foot with daily use ■ relieves itching, burning, cracking, and chafing associated with jock itch	
<b>Warnings</b>	
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<b>When using this product</b> ■ avoid spraying in eyes ■ use only as directed ■ intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal	
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<b>Other information</b> ■ store between 20° and 30°C (68° and 86°F) ■ if nozzle clogs, clean with a pin	
<b>Inactive ingredients</b> ISOBUTANE, SD ALCOHOL 40-B, KAOLIN, ZEA MAYS (CORN) STARCH, BHT, PPG-12-BUTETH-16	
<b>Questions? Call 1-866-964-0939</b>	



50-105QC-01

## TOLNAFATE

tolnaftate powder spray aerosol, spray

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:83324-234
<b>Route of Administration</b>	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
<b>ZEA MAYS (CORN) STARCH</b> (UNII: O8232NY3SJ)	
<b>BHT</b> (UNII: 1P9D0Z171K)	
<b>PPG-12-BUTETH-16</b> (UNII: 58CG7042J1)	
<b>ISOBUTANE</b> (UNII: BXR49TP611)	
<b>KAOLIN</b> (UNII: 24H4NWX5CO)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-234-46	130 g in 1 CAN; Type 0: Not a Combination Product	04/10/2025	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	04/10/2025	

**Labeler** - Chain Drug Marketing Association (011920774)

Revised: 11/2025

Chain Drug Marketing Association