

**TOLNAFATE- tolnaftate powder spray - talc free aerosol, spray
LEADER/ Cardinal Health 110, Inc.**

Leader Tolnaftate Antifungal 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)
- can prevent recurrence of most athlete's foot
- relieves symptoms of athlete's foot, including itching, burning, cracking, and chafing associated with jock itch

For external use only.

Flammable:

Do not use while smoking or near heat or flame. Do not puncture or incinerate. Contents under pressure. Do not store at temperature above 120°F. Intentional misuse by deliberately concentrating and inhaling contents can be harmful or fatal.

When using this product

- avoid contact with the eyes or mouth
- use only as directed

Stop use and ask a doctor if

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

Do not use

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

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 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
- use daily for 4 weeks for athlete's foot and ringworm; use daily for 2 weeks for jock itch
- if condition persists, consult a doctor
- this product is not effective on the scalp or nails
- if nozzle clogs, clean with a pin

Other information

store between 20° and 30°C (68°F and 86°)

Inactive ingredients

BHT, isobutane, kaolin, PPG-12-buteth-16, SD alcohol 40-B, zea mays (corn) starch

Questions?

Call 1-866-964-0939

Principal Display Panel

LEADER

Antifungal

Athlete's Foot

Powder Spray

Tolnaftate 1 % | Antifungal

Cure Most Athlete's Foot
and Prevents Recurrences
Relieves Itching, Cracking,
and Burning

Talc- Free

4.6 OZ (130 g)

Drug Facts

Active ingredient	Purpose
Tolnaftate 1%	Antifungal

Uses

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Warnings

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When using this product

- avoid contact with the eyes or mouth
- use only as directed

Stop use and ask a doctor if

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

Do not use on children under 2 years of age unless directed by a doctor.

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 over affected area twice daily (morning and night) or as directed by a doctor
- supervise children in the use of this product
- to prevent athlete's foot, apply once or twice daily (morning and/or night)
- 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
- use daily for 4 weeks for athlete's foot and ringworm; use daily for 2 weeks for jock itch
- if condition persists, consult a doctor
- this product is not effective on the scalp or nails
- if nozzle clogs, clean with a pin

Other information
store between 20° and 30° C (68° and 86° F)

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TOLNAFATE

tolnaftate powder spray - talc free aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0322
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	1.3 g in 130 g

Inactive Ingredients

Ingredient Name	Strength
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ISOBUTANE (UNII: BXR49TP611)	
KAOLIN (UNII: 24H4NWX5CO)	
PPG-12-BUTETH-16 (UNII: 58CG7042J1)	
ALCOHOL (UNII: 3K9958V90M)	

ZEA MAYS SUBSP. MAYS WHOLE (UNII: 1G5HNE09V8)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0322-1	130 g in 1 CAN; Type 0: Not a Combination Product	10/20/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	10/20/2017	

Labeler - LEADER/ Cardinal Health 110, Inc. (063997360)

Revised: 12/2025

LEADER/ Cardinal Health 110, Inc.