

TOLNAFTATE- jock itch powder aerosol, spray
Rite Aid Corporation

Rite Aid Antifungal Jock Itch Powder Spray Tolnaftate 1%

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- cures most jock itch (tinea cruris)
- relieves itching, burning and chafing associated with jock itch

Warnings

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 above 120°F. Intentional misuse by deliberately concentrating and inhaling contents can be harmful or fatal.

When using this product

- avoid contact with eyes
- use only as directed

Stop use and ask a doctor if

- irritation occurs
- no improvement within 2 weeks

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)

- supervise children in the use of this product
- use daily for 2 weeks; if conditions persists, consult a doctor
- if nozzle clogs, clean with a pin

Other information

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

Inactive ingredient

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

Questions?

Call 1-866-964-0939

Principal Display Panel

Compare to the active
ingredient, in **Tinactin**

Jock Itch Powder Spray

ANTIFUNGAL

JOCK ITCH

POWDER SPRAY

tolnaftate 1%

Cures most jock itch
Relieves itching, chafing & burning
Talc-free

NET WT

4.6 OZ (130 g)



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

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DISTRIBUTED BY: RITE AID
 30 HUNTER LANE, CAMP HILL, PA 17011
www.riteaid.com
 MADE IN USA WITH U.S. AND IMPORTED PARTS

SATISFACTION GUARANTEE:
 If you're not satisfied,
 we'll happily refund your money.



50-174RA

TOLNAFTATE

jock itch powder aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-0782
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
KAOLIN (UNII: 24H4NWX5CO)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ISOBUTANE (UNII: BXR49TP611)	

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:11822-0782-4	130 g in 1 CAN; Type 0: Not a Combination Product	03/15/2021	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M005	03/15/2021	

Labeler - Rite Aid Corporation (014578892)