

TOLNAFATE- tolnaftate jock itch powder spray - talc free aerosol, spray
Chain Drug Consortium, LLC

Premier Value Tolnaftate Jock Itch Powder Spray - Talc Free

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- cures most jock itch (tinea cruris)
- relieves itching, burning and crawling 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

When using this product

- do not get into eyes or mouth, if products get into eyes, rinse eyes thoroughly with water
- use only as directed

Intentional misuse by deliberately concentrating and inhaling contents can be harmful or fatal.

Stop use and ask a doctor if

- irritation occurs
- no improvement within 2 weeks

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away. Do not use on children under 2 years of age unless directed by a doctor.

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 persist, consult a doctor

- if nozzle clogs, clean with a pin

Other information

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

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**Premier Value****Antifungal****Jock Itch Powder Spray**

Tolnaftate 1%

Cures most jock itch

Relieves itching, chafing and burning



**Antifungal
Jock Itch
Powder Spray**

Tolnaftate 1%

Cures most jock itch

Relieves itching, chafing
and burning

Talc-free

NET WT 4.6 OZ (130 g)



COMPARE TO THE ACTIVE INGREDIENT IN TINACTIN®

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

Drug Facts	
Active ingredient	Purpose
Tolnaftate 1%	Antifungal
Uses	
<ul style="list-style-type: none"> ■ cures most jock itch (tinea cruris) ■ relieves itching, burning and chafing associated with jock itch 	
Warnings	
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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	
<ul style="list-style-type: none"> ■ use only as directed 	<ul style="list-style-type: none"> ■ avoid contact with eyes
Stop use and ask a doctor if	
<ul style="list-style-type: none"> ■ no improvement within 2 weeks 	<ul style="list-style-type: none"> ■ irritation occurs
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BHT, isobutane, kaolin, PPG-12-buteth-16, SD alcohol 40-B, zea mays (corn) starch	
Questions? Call 1-866-964-0939	

Distributed By:
Pharmacy Value Alliance, LLC
407 East Lancaster Avenue
Wayne, PA 19087
Made in USA with U.S.
and imported parts



If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.

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FPO

50-061 PV

Talc- Free

NET WT 4.6 OZ (130 g)

TOLNAFATE

tolnaftate jock itch powder spray - talc free aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-653
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:68016-653-46	130 g in 1 CAN; Type 0: Not a Combination Product	10/31/2017	06/30/2028

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	10/31/2017	06/30/2028

Labeler - Chain Drug Consortium, LLC (101668460)

Revised: 2/2026

Chain Drug Consortium, LLC