

TINACTIN JOCK ITCH- tolnaftate aerosol, powder
Bayer HealthCare LLC

Tinactin Jock Itch Powder Spray Talc Free

Drug Facts

Active ingredient

(To Deliver) Tolnaftate 1%

Purpose

Antifungal

Uses

- cures most jock itch
- for effective relief of itching, chafing and burning

Warnings

For external use only

Flammable: Do not use near heat, flame, or while smoking

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

When using this product

- avoid contact with the eyes
- use only as directed. Intentional misuse by deliberately concentrating and inhaling contents can be harmful or fatal
- contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120°F.

Stop use and ask a doctor if

- irritation occurs
- there is no improvement within 2 weeks

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 condition persists longer, ask a doctor

- this product is not effective on the scalp or nails

Other information

store between 20° to 25°C (68° to 77°F)

Inactive ingredients

butylated hydroxytoluene, hydroxypropyl cellulose, isobutane, kaolin, magnesium stearate, PPG-12-buteth-16, SD alcohol 40-B (9% w/w), zea mays (corn) starch

Questions?

1-866-360-3266 or visit us at www.tinactin.com

PRINCIPAL DISPLAY PANEL - 133g Can Label

TOUGH ACTIN' ®

Tinactin®

tolnaftate **ANTIFUNGAL**

Cures Most Jock Itch

Relieves:

- Itching
- Burning

POWDER SPRAY

Goes On Dry

Talc Free

NET WT 133g (4.6 oz)



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tolnaftate ANTIFUNGAL

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Questions? 1-866-360-3296 or visit us at www.tinactin.com

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Dist. by: Bayer HealthCare LLC
Whispery, NJ 07981

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TINACTIN JOCK ITCH

tolnaftate aerosol, powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11523-0138
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
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ISOBUTANE (UNII: BXR49TP611)	
PPG-12-BUTETH-16 (UNII: 58CG7042J1)	

KAOLIN (UNII: 24H4NWX5CO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white (white to off-white)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-0138-1	133 g in 1 CAN; Type 0: Not a Combination Product	03/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	03/01/2024	

Labeler - Bayer HealthCare LLC (112117283)

Revised: 4/2024

Bayer HealthCare LLC