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 cruis)
- relieves itching, burning and crafing 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 120ºF

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 cans 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 20º and 30ºC (68ºF and 86ºF)

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

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