TOLNAFTATE- antifungal spray liquid aerosol, spray
Chain Drug Consortium, LLC

Premier Value Antifungal Spray Liquid

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- proven clinically effective in the treatment of athlete's foot (tinea pedis) and ringworm (tinea corporis)
- prevents the recurrence of most athlete's foot with daily use
- for effective relief of itching, cracking and burning

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 at temperature above 120ºF. 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

- irritation occurs
- no improvement within 4 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

- 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
- if condition persists, consult a doctor
- to prevent athlete's foot, apply once or twice daily (morning and/or night)
- this product is not effective on the scalp or nails
- if nozzle clogs, clear under running water

Other information

store between 20º and 30ºC (68º and 86ºF)

Inactive ingredients

BHT, isobutane, PPG-12-buteth-16, SD alcohol 40-B

Questions?

Call 1-866-964-0939

Principal Display Panel

Premier Value

Antifungal

Athlete's Foot

Liquid Spray

Tolnaftate 1%

Cures most athlete's foot and prevents recurrences

Relieves itching, cracking and burning

Net WT 5.3 OZ (150g)

TOLNAFTATE

antifungal spray liquid aerosol, spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-609
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	0.15 g in 150 g

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ISOBUTANE (UNII: BXR49TP611)	
PPG-12-BUTETH-16 (UNII: 58CG7042J1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-609-00	150 g in 1 CAN; Type 0: Not a Combination Product	05/23/2003	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	05/23/2003	

Labeler - Chain Drug Consortium, LLC (101668460)

Revised: 12/2024 Chain Drug Consortium, LLC