

TOLNAFTATE- tolnaftate aerosol, spray

Target Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Antifungal Liquid Spray

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- cures most athlete's foot (tinea pedis) and ringworm (tinea corporis)
- if used daily, can prevent recurrence of athlete's foot
- relieves symptoms of athlete's foot, including itching, burning and cracking

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 120°F.

When using this product

- do not get into eyes or mouth
- 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 four weeks (for athlete's foot) or two weeks (for jock itch)

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 two 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) or as directed by a doctor
- 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 conditions persist, consult a doctor
- to prevent athlete's foot, appl morning and/or night
- this product is not effective on the scalp or nails

▯ ***Other information***

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

▯ ***Inactive ingredients***

BHT, isobutane (propellant), PPG-12-buteth-16, SD alcohol 40-B(36%w/w)

▯ ***Questions?***

call 1-800-910-6874

Principal Display Panel

antifungal liquid spray

tolnaftate 1%

Compare to Tinactin® Liquid Spray

cures and prevents most athlete's foot

cures most jock itch and ringworm

helps relieve burning and itching

up&up

Net WT 5.3 OZ (150.2g)▯

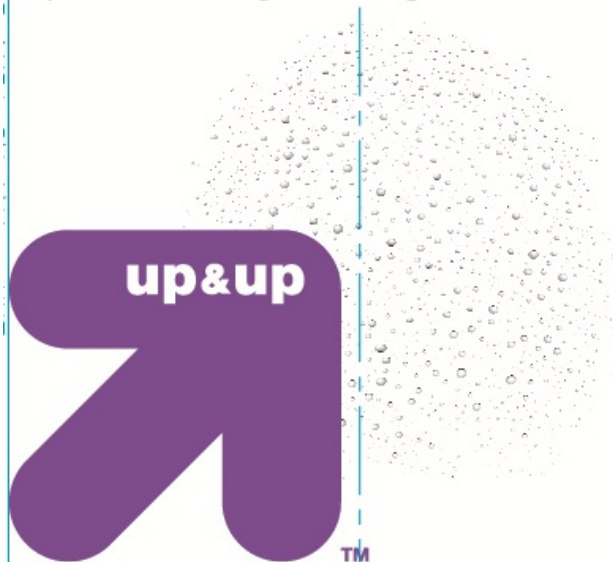
antifungal liquid spray

NDC 11673-448*5

tolnaftate 1%

Compare to Tinactin® Liquid Spray*

cures and prevents most athlete's foot
cures most jock itch and ringworm
helps relieve burning and itching



NET WT 5.3 OZ (150.2 g)

Drug Facts

Active ingredient	Purpose
Tolnaftate 1%	Antifungal

Uses

- cures most athlete's foot (tinea pedis), ringworm (tinea corporis) and jock itch (tinea cruris)
- if used daily, can prevent recurrence of athlete's foot
- relieves symptoms of athlete's foot, including itching, burning and cracking

Warnings

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- supervise children in the use of this product
- for athlete's foot, pay special attention to spaces between 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 daily (morning and/or night)
- this product is not effective on scalp or nails

Other information

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

Inactive ingredients

BHT, isobutane (propellant), PPG-12-buteth-16, SD alcohol 40-B (36% w/w)

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245 06 0405 ID321468

Distributed by Target Corp., Mpls, MN 55403

Made in U.S.A.

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ALTSTLC-1



TOLNAFTATE

antifungal liquid spray aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG LABEL	Item Code (Source)	NDC:11673-448
Route of Administration	TOPICAL	DEA Schedule	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TOLNAFTATE (TOLNAFTATE)	TOLNAFTATE	1.5 g in 150.2 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL	
BUTYLATED HYDROXYTOLUENE	
ISOBUTANE	
PPG-12-BUTETH-16	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-448-05	150.2 g in 1 CANISTER		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333C	12/31/2008	

Labeler - Target Corporation (006961700)**Establishment**

Name	Address	ID/FEI	Business Operations
Premier Brands of America Inc.		063849780	relabel(11673-448)

Revised: 9/2013

Target Corporation