

**ANTIFUNGAL LIQUID- tolnaftate liquid**  
**Target Corporation**

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**UP & UP Anti-Fungal Liquid**

***Active ingredient***

Tolnaftate 1%

***Purpose***

Antifungal

***Uses***

proven clinically effective in the treatment of most 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, burning, cracking, redness and scaling

***Warnings***

**For external use only.**

**Do not use**

on children under 2 years of age unless directed by a doctor.

**When using this product**

avoid contact with the eyes.

**Stop use and consult a doctor if**

- irritation occurs
- there is no improvement within 4 weeks

**Keep out of reach of children.**

If accidental ingestion occurs, get medical help or contact a Poison Control Center right away.

***Directions***

- clean the affected area and dry thoroughly
- apply a thin layer of the product over the 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
- for athlete's foot and ringworm, use daily for 4 weeks.
- if condition persists longer, consult a doctor
- to prevent athlete's foot: wash the feet and dry thoroughly; apply a thin layer of the product to the feet once or twice daily (morning and/or night)
- this product is not effective on the scalp or nails

### ***Other information***

- SHAKE WELL BEFORE USE
- store at room temperature 15°-30°C (59° - 86°F)
- protect from freezing; if freezing occurs, warm to room temperature
- keep tightly closed when not in use

### ***Inactive ingredient***

WATER, GLYCERIN, SCLEROTIUM GUM, PHENOXYETHANOL, DISODIUM EDTA, ETHYLHEXYLGLYCERIN, XANTHAN GUM

### ***Questions?***

**Call 1-866-964-0939**

### **Principal Display Panel**

MAXIMUM STRENGTH

Antifungal Liquid

TOLNAFTATE 1%/ ANTIFUNGAL

Proven clinically effective in

the treatment of fungus on fingers, toes and skin around nails

Prevents the recurrence

of most athlete's foot

with daily use

Relieves Itching, Burning

& Scaling

Built-in Brush for Easy

Application

With moisturizing Ingredients

SHAKE WELL BEFORE USE

NET 1 FL. OZ. (30 mL)



**Maximum Strength  
Antifungal  
Liquid**

Tolnaftate 1% / Antifungal



**DO NOT USE IF NECKBAND IS BROKEN**

**Drug Facts**

**Active ingredient Purpose**  
Tolnaftate 1% Antifungal

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up&up™ Maximum Strength Antifungal Liquid treats and helps stop the spread of fungal infections on cuticles, around nail edges, and skin under nail tips where reachable with applicator brush. Topical antifungal treatments will not penetrate hard nail surfaces.

**SHAKE WELL BEFORE USE**

\*This product is not manufactured or distributed by Kramer Laboratories, Inc., owner of the registered trademark Fungi-Nail®.

**Satisfaction guaranteed – Love it or your money back.**

Distributed by Target Corporation  
Minneapolis, MN 55403  
Made in Canada  
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Compare to active ingredient in Fungi-Nail® Anti-Fungal Liquid\*

**Maximum Strength  
Antifungal  
Liquid**

Tolnaftate 1% / Antifungal

- Proven clinically effective in the treatment of Fungus on fingers, toes and skin around nails
- Prevents the recurrence of most athlete's foot with daily use
- Relieves itching, burning and scaling
- Built-in brush for easy application
- With moisturizing ingredients
- SHAKE WELL BEFORE USE



Easy to Apply  
Applicator  
Brush



1 FL. OZ. (30 mL)



Actual Size

53-121TG-02

**ANTIFUNGAL LIQUID**

tolnaftate liquid

**Product Information**

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:82442-956

**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 mL

### Inactive Ingredients

Ingredient Name	Strength
XANTHAN GUM (UNII: TTV12P4NEE)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BETASIZOFIRAN (UNII: 2X51AD1X3T)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-956-01	1 in 1 CARTON	07/19/2024	11/30/2026
1		30 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	07/19/2024	11/30/2026

**Labeler** - Target Corporation (006961700)

Revised: 8/2025

Target Corporation