

ANTIFUNGAL LIQUID- tolnaftate liquid
Target Corporation

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 NECKLAD IS BROKEN

Drug Facts

Active ingredient Purpose
Tolnaftate 1% Antifungal

Uses

- proven clinically effective in the treatment of most athlete's foot (tinea pedis) and ringworm (tinea corporis)
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Questions? Call 1-866-964-0839

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.

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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

DO NOT USE IF NECKBAND IS BROKEN

Active ingredient: Tolnaftate (1%)

Purpose: Antifungal **Uses:** Proven clinically effective in the treatment of most athlete's foot (tinea pedis) and ringworm (tinea corporis)

Maximum Strength Antifungal Liquid

Tolnaftate 1% / Antifungal

- Proven clinically effective in the treatment of fungus on fingers, toes and skin around nails
- SHAKE WELL BEFORE USE



1 FL. OZ. (30 mL)

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 ask a doctor if irritation occurs or 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. **Inactive Ingredients:** WATER, GLYCERIN, SCLEROTIUM GUM, PHENOXYETHANOL, DISODIUM EDTA, ETHYLHEXYLGLYCERIN, XANTHAN GUM.

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. Keep packaging for complete drug facts information.

Questions? Call 1-866-964-0939

245 06 0098 R00 C-002262-01-098

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50-121TG-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
GLYCERIN (UNII: PDC6A3C0OX)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BETASIZOFIRAN (UNII: 2X51AD1X3T)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:82442-956-01	1 in 1 CARTON	07/19/2024	
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	

Labeler - Target Corporation (006961700)

Revised: 7/2024

Target Corporation