

UP AND UP ANTIFUNGAL- tolnaftate 1% cream
Target Corporation

Up and Up Antifungal Cream

DRUG FACTS

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- for effective treatment of most athlete's foot (tinea pedis), jock itch (tinea cruris) and ringworm (tinea corporis)
- for effective relief of itchy, scaly skin between the toes
- clears up most athlete's foot infection and with daily use helps keep it from coming back

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 eyes

Stop use and ask a doctor if

- irritation occurs or if there is no improvement within 4 weeks (for athlete's foot and ringworm)
- irritation occurs or if there is no improvement within 2 weeks (for jock itch)

Keep this and all drugs out of the reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

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

- use daily for 4 weeks. If condition persists longer, consult a doctor
- pay special attention to the spaces between the toes
- wear well fitting ventilated shoes
- change shoes and socks at least once daily

For ringworm, use daily for 4 weeks. If condition persists longer, consult a doctor.
For jock itch, use daily for 2 weeks. If condition persists longer, consult a doctor.
This product is not effective on the scalp or nails.

Other information

store at controlled room temperature 20° -25°C (68-77°F)

Inactive ingredients

Ceteth-20, Cetostearyl Alcohol, Chlorocresol, Mineral Oil, Petrolatum , Propylene Glycol, Sodium Phosphate Monobasic, Water (Purified)

Questions or comments? call 1-800-910-6874

Principal display panel

Up &Up NDC 82442-020-28

Antifungal Cream

Tolnaftate 1%

NET WT 1 OZ (28g)

Compare to active ingredient in Tinactin®*

Antifungal Cream



up&up™

Tolnaftate 1%

- Cures and prevents most athlete's foot
- Helps relieve itching and burning

NET WT 1 OZ (28 g)

LOT
EXP

Satisfaction guaranteed –
Love it or your money back.

245 06 0415 R00
C-002262-01-098-0000
Distribution by Target Corporation
Minneapolis, MN 55403
Made in U.S.A. with U.S. and imported
ingredients and components
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NDC 80442-000-28



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*This product is not manufactured or distributed by Bayer Corporation, owner of the registered trademark, Tinactin™.

Drug Facts	Drug Facts (continued)
Active ingredients Tolnaftate 1% Antifungal	
Purpose Antifungal	
Uses • proven clinically effective in the treatment of most athlete's foot • helps prevent most athlete's foot with daily use • helps relieve most athlete's foot with daily use	
Warnings For external use only Do not use on children under 2 years of age except under the advice and supervision of a doctor. When using this product avoid contact with the eyes. Stop use and ask a doctor if • there is no improvement within 4 weeks • irritation occurs • there is no improvement within 4 weeks Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away at (1-800-222-1222).	
Directions • wash affected area and dry thoroughly • apply 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 toes, wear wet-drying, ventilated shoes and change shoes and socks at least once daily • use daily for 4 weeks; if condition persists longer, ask 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	
Other information • Store at controlled room temperature 20°–25°C (68°–77°F)	
Inactive ingredients Cetyl-20, Cetylstearyl Alcohol, Cholesterol, Mineral Oil, Petroleum, Propylene Glycol, Sodium Phosphate Monobasic, Water (Purified)	
Questions or comments? Call 1-800-910-6874	

up&up



Antifungal Cream
up&up Tolnaftate 1%
 • Cures and prevents most athlete's foot
 • Helps relieve itching and burning
 NET WT 1 OZ (28 g)

Active ingredient: Tolnaftate 1% **Purpose:** Antifungal
Uses: • proven clinically effective in the treatment of most athlete's foot (tinea pedis) and ringworm (tinea corporis)
 • helps prevent most athlete's foot with daily use • for effective relief of itching, burning and cracking
Warnings: for external use only. Do not use on children under 2 years of age except under the advice and supervision of a doctor
When using this product avoid contact with the eyes Stop use and ask a doctor if • irritation occurs
 • there is no improvement within 4 weeks
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away at (1-800-222-1222).
Directions: • wash affected area and dry thoroughly • apply 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 longer, ask 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
Questions? Call 1-800-910-6874

NDC 82442-020-28
 245-061015 R00
 C-002262-01-098-0000
 Dist. by Target Corp.
 Mpls., MN 55403
Made in U.S.A.
 with U.S. and imported ingredients and components
 TM & ©2024 Target Brands, Inc. 201060

UP AND UP ANTIFUNGAL

tolnaftate 1% cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82442-020
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 g

Inactive Ingredients

Ingredient Name	Strength
CETETH-20 (UNII: I835H2IHHX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CHLOROCRESOL (UNII: 36W53O7109)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPUU)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-020-28	1 in 1 CARTON	08/01/2024	

1	28 g in 1 TUBE; 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	08/01/2024	

Labeler - Target Corporation (006961700)

Registrant - Sheffield Pharmaceuticals LLC (151177797)

Establishment			
Name	Address	ID/FEI	Business Operations
Sheffield Pharmaceuticals LLC		151177797	manufacture(82442-020) , analysis(82442-020)

Revised: 5/2024

Target Corporation