

ATHLETES FOOT- butenafine hydrochloride cream

Target Corporation

Athlete's Foot Cream

Drug Facts

Active ingredient

Butenafine hydrochloride 1%

Purpose

Antifungal

Uses

- cures most athlete's foot between the toes.
Effectiveness on the bottom or sides of foot is unknown.
- cures most jock itch and ringworm
- relieves itching, burning, cracking, and scaling which accompany these conditions

Warnings

For external use only

Do not use

- on nails or scalp
- in or near the mouth or the eyes
- for vaginal yeast infections

When using this product do not get into the eyes. If eye contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if too much irritation occurs or irritation gets worse

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- adults and children 12 years and older:
 - use the tip of the cap to break the seal and open the tube
 - wash the affected skin with soap and water and dry completely before applying
 - **for athlete's foot between the toes:** apply to affected skin between and around the toes twice a day for 1 week (morning and night), or once a day for 4 weeks, or as directed by a doctor. Wear well-fitting, ventilated shoes. Change shoes and socks at least once daily.

Apply between and around the toes



1 week twice a day or 4 weeks once a day

- **for jock itch and ringworm:** apply once a day to affected skin for 2 weeks or as directed by a doctor.

- wash hands after each use
- children under 12 years: ask a doctor

Other information

- do not use if seal on tube is broken or not visible
- store between 20° to 25° C (68° to 77° F)

Inactive ingredients

benzyl alcohol, cetyl alcohol, glycerin, glyceryl monostearate SE, polyoxyethylene (23) cetyl ether, propylene glycol dicaprylate, purified water, sodium benzoate, stearic acid, trolamine, white petrolatum

Questions?

Call **1-800-910-6874**

Distributed by Target Corporation
Minneapolis, MN 55403

PRINCIPAL DISPLAY PANEL - 30 g Tube Carton

up&up

ultra

antifungal cream

butenafine hydrochloride cream 1%

clinically proven to cure most athlete's foot between toes
prescription strength
relieves itching, burning and cracking

1

WEEK

TREATMENT**

****option for**

athlete's foot.

See directions.

NET WT 1 OZ (30 g)

Contains the Drug BUTENAFINE HYDROCHLORIDE

NDC 11673-016-02



Compare to active ingredient in Lotrimin Ultra®

ultra antifungal cream

butenafine hydrochloride cream 1%

1
WEEK
TREATMENT**

**option for athlete's foot. See directions.

LPK-8680-0
1217-0
52



T175
B76.2
ENG19.54



ultra antifungal cream

butenafine hydrochloride cream 1%

clinically proven to cure most athlete's foot between toes prescription strength relieves itching, burning and cracking

NET WT 1 OZ (30 g)

1
WEEK
TREATMENT**

**option for athlete's foot. See directions.

NO VARNISH/NO AD
NO COPY/NO COLOR
THIS FLAP FOR LOT #
AND EXP DATE PRINT

245 06 0403 R00
C-000703-01-028



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or your money back.

We welcome any questions you may have at
Target.com/comments or 1-800-910-6874.

Each year we give 5% of our profit to communities. See
all the good we do together at Target.com/Community.

*All trademarks are property of their respective owners. This product is
not affiliated with the makers/owners of Lotrimin Ultra®.

NO VARNISH
ON THIS FLAP

Drug Facts	
Active ingredient Butenafine hydrochloride 1% Antifungal	
Uses • cures most athlete's foot between the toes. • Electrolytes on the bottom or sides of foot is unknown. • cures most foot itch and ringworm. • relieves itching, burning, cracking, and scaling which accompany these conditions.	
Warnings For external use only. Do not use: • on nails or scalp • in or near the mouth or the eyes • for vaginal yeast infections When using this product do not get it in the eyes. If eye contact occurs, rinse thoroughly with water. Stop use and ask a doctor if too much irritation occurs or irritation gets worse. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions • adults and children 12 years and older: • use the tip of the cap to break the seal and open the tube • wash the affected skin with soap and water and dry completely before applying • for the foot: apply to affected skin between and around the toes twice a day for 1 week (from ingrown nail), or once a day for 4 weeks, or as directed by a doctor. Wear well-fitting, ventilated shoes. Change shoes and socks at least once daily. • for foot, hand and groin rashes: • apply once a day to affected skin • apply between mandatory soaks • wash hands after each use • children under 12 years: ask a doctor. 1 week, 4 weeks, or 7 weeks, as directed.	
Other information • do not use if seal on tube is broken or not visible • store between 20° to 25° C (68° to 77° F) Inactive ingredients benzyl alcohol, cetyl alcohol, glycerin, glyceryl monostearate SE, polyoxyethylene (20) cetyl ether, propylene glycol decapylate, purified water, sodium benzoate, stearic acid, tolnamine, white petrolatum	
Questions? Call 1-800-910-6874	

NO VARNISH
ON THIS FLAP

ATHLETES FOOT

butenafine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-016
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Butenafine Hydrochloride (UNII: R8 XA2029 ZI) (Butenafine - UNII:9 1Y494NL0 X)	Butenafine Hydrochloride	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
benzyl alcohol (UNII: LKG8494WBH)	
cetyl alcohol (UNII: 936JST6JCN)	
glycerin (UNII: PDC6A3C0OX)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
CETETH-23 (UNII: 495CTZ441V)	
propylene glycol dicaprylate (UNII: 581437HWX2)	
WATER (UNII: 059QF0KO0R)	
sodium benzoate (UNII: OJ245FE5EU)	
stearic acid (UNII: 4ELV7Z65AP)	
trolamine (UNII: 9O3K93S3TK)	
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics

Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-016-02	1 in 1 CARTON	11/30/2017	
1		30 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
ANDA	ANDA205181	11/17/2017	

