

ATHLETES FOOT- butenafine hydrochloride cream
CVS Pharmacy

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 1-866-923-4914

Distributed by: CVS Pharmacy, Inc.

One CVS Drive, Woonsocket, RI 02895

PRINCIPAL DISPLAY PANEL - 30 g Tube Carton

CVSHealth™

**Butenafine Hydrochloride
Cream 1%**

**Antifungal Cream
Prescription Strength**

NET WT 1 OZ (30 g)

Package Contains One Tube



Compare to the active ingredient in Lotrimin Ultra®*

Prescription Strength:

- Relieves itching, burning, and cracking
- Clinically proven to cure most athlete's foot between the toes

Contains the Drug:
BUTENAFINE HYDROCHLORIDE

1 Week
Treatment Option
For Athlete's Foot
(See Directions)



Butenafine Hydrochloride Cream 1%

Antifungal Cream Prescription Strength

LPK-8409-1
1117-1
611

T175
B76.2
ENG19.54



Butenafine Hydrochloride Cream 1%

Antifungal Cream Prescription Strength

NET WT 1 OZ (30 g)

Package Contains One Tube



Butenafine Hydrochloride Cream 1%

Antifungal Cream Prescription Strength

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

#270832



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Made in Canada
V-11653



NO VARNISH
ON THIS FLAP

Drug Facts

Active ingredient
Butenafine hydrochloride 1% Antifungal

Purpose
Antifungal

Uses
• cures most athlete's foot between the toes.
Effectiveness on the bottom or sides of foot is unknown.
• cures most itchy, burning, cracking, and scaling which accompanies these conditions

Warnings
For external use only
Do not use
• on or near the mouth or the eyes
• in or near the mouth or the eyes
• if you have eye 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.

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 (28) cetyl ether, propylene glycol, decylate, purified water, sodium benzoate, stearic acid, toluene, white petrolatum

Directions
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Apply between and around the toes.
• for itchy, burning, cracking, and scaling which accompanies these conditions:
• 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 if you have any eye or skin infections

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 (28) cetyl ether, propylene glycol, decylate, purified water, sodium benzoate, stearic acid, toluene, white petrolatum

Questions? Call 1-866-923-4914

Contains the Drug: BUTENAFINE HYDROCHLORIDE



butenafine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-986
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Butenafine Hydrochloride (UNII: R8 XA2029ZI) (Butenafine - UNII:9 1Y494NL0X)	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:69842-986-08	1 in 1 CARTON	11/17/2017	
1		12 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69842-986-02	1 in 1 CARTON	11/17/2017	
2		30 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:69842-986-03	1 in 1 CARTON	11/21/2019	
3		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	

Labeler - CVS Pharmacy (062312574)

Revised: 1/2019

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