

ATHLETES FOOT ULTRA- butenafine hydrochloride cream H E B

Athlete's Foot Cream Ultra

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**

PRINCIPAL DISPLAY PANEL - 12 g Tube Carton

Compare to Lotrimin Ultra [®]* active ingredient

NDC 37808-013-02

H-E-B [®]

Clinically Proven to Cure Most Athlete's Foot between the Toes

Athlete's Foot Cream Ultra

Butenafine Hydrochloride Cream 1%

Antifungal

NET WT 12 g (0.42 oz)



NDC 37808-013-02

Clinically Proven to Cure Most Athlete's Foot between the Toes

Athlete's Foot Cream Ultra

Butenafine Hydrochloride Cream 1%

Antifungal

Contains the Drug: BUTENAFINE HYDROCHLORIDE

1 Week

Treatment Option For Athlete's Foot - See Directions

T174
B75.2
ENG19.53

- Prescription Strength
- Relieves Itching, Burning and Cracking

Compare to Lotrimin Ultra[®] active ingredient



NDC 37808-013-02

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Athlete's Foot Cream Ultra

Butenafine Hydrochloride Cream 1%

Antifungal

NET WT 12 g (0.42 oz)

Athlete's Foot Cream Ultra

Butenafine Hydrochloride Cream 1%

Antifungal

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

LPK-8788-1
0818-1
58

100% GUARANTEE promise

If you aren't completely pleased with this product, we'll be happy to replace it or refund your money. You have our word on it.

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



MADE WITH PRIDE & CARE FOR H-E-B[®]
SAN ANTONIO, TX 78204
MADE IN CANADA

NO VARNISH ON THIS FLAP

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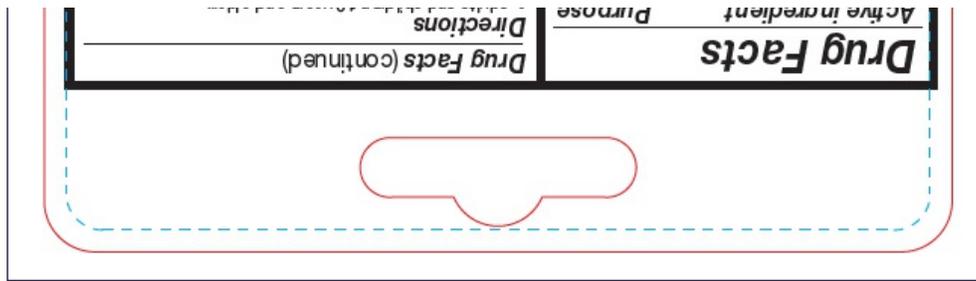
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Apply between socks at least once daily.

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- 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 4 weeks once a day

1 week 1 a day or 4 weeks once a day



ATHLETES FOOT ULTRA

butenafine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-013
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BUTENAFINE HYDROCHLORIDE (UNII: R8XA2029ZI) (BUTENAFINE - UNII:91Y494NLOX)	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:37808-013-02	1 in 1 CARTON	12/12/2017	
1		12 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 - H E B (007924756)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(37808-013)

Revised: 7/2025

H E B