

BUTENAFINE HYDROCHLORIDE- butenafine hydrochloride cream
YYBA CORP

Butenafine Hydrochloride

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 **1-866-933-6337**

Distributed by:

Wellspring

Airmont, NY 10952, U.S.A.

PRINCIPAL DISPLAY PANEL - 30 g Tube Carton

Butenafine

Hydrochloride Cream 1%

Antifungal

NET WT 30 g (1 oz)



Compare to the active ingredient in Lotrimin Ultra^{®*}

Clinically Proven to Cure Most ATHLETE'S FOOT Between the Toes

Contains the Drug: BUTENAFINE HYDROCHLORIDE

Relieves: Itching Burning Cracking

1 Week
Treatment Option
For Athlete's Foot
See Directions

Prescription Strength

52A2625
0823
11

T175
B76.2
ENG19.54

Butenafine

Hydrochloride Cream 1%

Antifungal

NET WT 30 g (1 oz)

Butenafine
Hydrochloride
Cream 1%
Antifungal

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



3 73581 00025 8

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

Distributed by:
Wellspring
Aimont, NY 10952, U.S.A.



Why pay more?
wellspringmeds.com
866-933-6337

Made in Canada.

NO VARNISH
ON THIS FLAP

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Butenafine
Hydrochloride Cream 1%
Antifungal
Clinically Proven to Cure Most Athlete's Foot Between the Toes
Contains the Drug: BUTENAFINE HYDROCHLORIDE

BUTENAFINE HYDROCHLORIDE

butenafine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73581-210
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73581-210-02	1 in 1 CARTON	10/12/2023	
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	10/12/2023	

Labeler - YYBA CORP (006339772)

Registrant - Taro Pharmaceuticals U.S.A., Inc. (145186370)

Establishment

Name	Address	ID/FEI	Business Operations
Taro Pharmaceuticals Inc.		206263295	MANUFACTURE(73581-210)

Revised: 10/2023

YYBA CORP