JOCK ITCH CREAM- butenafine hydrochloride cream
TARGET CORPORATION

Target Butenafine HCl Cream 1%- 12 g

Drug Facts

Butenafine Hydrochloride 1%

Antifungal

Uses

- cures most jock itch
- relieves itching, burning, crackling, and scaling which accompany this condition

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 irration gets worse.

Keep out of reach of children. If swallowed, get medical help or contact a Poision 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
- 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)

Benzyl Alcohol, Cetyl Alcohol, Glycerin, Glyceryl Monostearate SE, Polyoxyethylene (23)

Cetyl Ether, Propylene Glycol Dicaprylate, Purified Water, Sodium Benzoate, Stearic Acid, Trolamine, White Petrolatum.

Question?

1-800-910-6874

DISTRIBUTED BY:

Target Corporation.,

Minneapolis, MN 55403

PRINCIPAL DISPLAY PANEL - 12 g Tube Carton

Compare to active ingredient in Lotrimin Ultra ®Jock Itch*

Jock Itch

Cream

Butenafine Hydrochloride

Cream 1%

- Prescription strength
- Relieves itching, burning and chafing
- Clinically proven to cure most jock itch
- Contains the drug:

butenafine hydrochloride

up&up ™

NET WT 0.42 oz (12 g)

JOCK ITCH CREAM

butenafine hydrochloride cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-436
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BUTENAFINE HYDROCHLORIDE (UNII: R8XA2029ZI) (BUTENAFINE - UNII:91Y494NL0X)	BUTENAFINE HYDROCHLORIDE	10 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETETH-23 (UNII: 495CTZ441V)	
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TROLAMINE (UNII: 9O3K93S3TK)	
PROPYLENE GLYCOL DICAPRYLATE (UNII: 581437HWX2)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	white	**Score**	

Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-436-01	1 in 1 CARTON	11/15/2024	
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	11/15/2024	

Labeler - TARGET CORPORATION (006961700)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(11673-436)

Revised: 7/2025 TARGET CORPORATION