UP AND UP JOCK ITCH RELIEF- clotrimazole cream
Target Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Target Corporation Jock Itch Relief Drug Facts

Active ingredient

Clotrimazole 1%

Purpose

Antifungal

Uses

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

Warnings

For external use only

Do not use

on children under 2 years of age unless directed by a doctor

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

- irritation occurs
- there is no improvement within 2 weeks

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- wash the affected area and dry thoroughly
- apply a thin layer over affected area twice daily (morning and night)
- supervise children in the use of this product
- use daily for 2 weeks
- if condition persists longer, ask a doctor
- this product is not effective on the scalp or nails

Other information

- store at 20°-25°C (68°-77°F)

Inactive ingredients

benzyl alcohol, cetostearyl alcohol, cetyl esters wax, octyldodecanol, polysorbate 60, purified water, sorbitan monostearate

Questions?

Call 1-888-547-7400

Package/Label Principal Display Panel

Compare to active ingredient in Lotrimin® AF

jock itch relief

antifungal cream

clotrimazole 1%

cures most jock itch

relieves itching, burning, scaling and chafing

NET WT 0.5 OZ (14 g)

UP AND UP JOCK ITCH RELIEF

clotrimazole cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-504
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)	CLOTRIMAZOLE	1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ESTERS WAX (UNII: D072FFP9GU)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
WATER (UNII: 059QF0KO0R)	
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
OCTYLDODECANOL (UNII: 461N1O614Y)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-504-58	1 in 1 CARTON	09/17/2013	
1		14 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
OTC monograph final	part333C	09/17/2013	

Labeler - Target Corporation (006961700)

Revised: 7/2016 Target Corporation