LOTRIMIN- clotrimazole cream Bayer HealthCare LLC

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.

Lotrimin ® Clotrimazole

Drug Facts

Active ingredient

Clotrimazole 1%

Purpose

Antifungal

Uses

- cures most athlete's foot, jock itch, and ringworm
- relieves itching, burning, cracking, scaling and discomfort which accompany these conditions

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 4 weeks (for athlete's foot and ringworm) or 2 weeks (for jock itch)

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- wash 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
- for athlete's foot: pay special attention to spaces between the toes, wear well-fitting, ventilated shoes and change shoes and socks at least once daily
- for athlete's foot and ringworm, use daily for 4 weeks; for jock itch, 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 between 20° to 25°C (68° to 77°F)

Inactive ingredients

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

Questions?

866-360-3226

Distributed by Bayer HealthCare LLC, Whippany, NJ, USA, 07981

PRINCIPAL DISPLAY PANEL - 12 g Tube Carton



LOTRIMIN [®]AF clotrimazole cream ANTIFUNGAL

NET WT 12g (0.42 OZ)

LOTRIMIN clotrimazole cream Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:11523-0963 Route of Administration TOPICAL Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)	CLOTRIMAZOLE	10 mg in 1 g

Inactive Ingredients			
Ingredient Name	Strength		
BENZYL ALCOHOL (UNII: LKG8494WBH)			
CETYL ALCOHOL (UNII: 936JST6JCN)			
CETYL ESTERS WAX (UNII: D072FFP9GU)			
OCTYLDODECANOL (UNII: 461N1O614Y)			
POLYSORBATE 60 (UNII: CAL22UVI4M)			
SORBITAN MONOSTEARATE (UNII: NVZ 4I0H58X)			
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)			
WATER (UNII: 059QF0KO0R)			

Product Characteristics			
Color	white (White to Off-white)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:11523- 0963-1	1 in 1 CARTON	02/01/2002			
1		30 g in 1 TUBE; Type 0: Not a Combination Product				
2	NDC:11523- 0963-2	1 in 1 CARTON	02/01/2002			
2		15 g in 1 TUBE; Type 0: Not a Combination Product				
3	NDC:11523- 0963-5	1 in 1 CARTON	02/01/2002			
3		12 g in 1 TUBE; Type 0: Not a Combination Product				
4	NDC:11523- 0963-7	1 in 1 CARTON	02/01/2002			
4		24 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	02/01/2002		

Revised: 9/2023 Bayer HealthCare LLC