CLOTRIMAZOLE- clotrimazole cream NuCare Pharmaceuticals, Inc.

Perrigo Clotrimazole Cream USP, 1% 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. (1-800-222-1222)

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
- for athlete's foot pay special attention to the spaces between the toes, wear wellfitting 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, consult a doctor
- this product is not effective on the scalp or nails

Other information

store at 20°-25°C (68°-77°F)

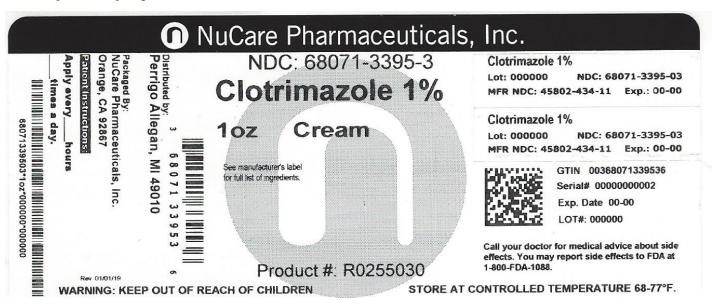
Inactive ingredients

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

Questions or comments?

1-800-719-9260

Principal Display Panel



CLOTRIMAZOLE clotrimazole cream Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:68071-3395(NDC:45802-434) 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)				
OCTYLDODECANOL (UNII: 461N1O614Y)				
POLYSORBATE 60 (UNII: CAL22UVI4M)				
WATER (UNII: 059QF0KO0R)				
SORBITAN MONOSTEARATE (UNII: NVZ 410H58X)				

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
NDC:68071- 3395-3	30 g in 1 BOX; Type 0: Not a Combination Product	07/12/2017			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M014	06/03/2011				

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment						
Name	Address	ID/FEI	Business Operations			
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-3395)			

Revised: 7/2024 NuCare Pharmaceuticals, Inc.