

CLOTRIMAZOLE CREAM 1%- clotrimazole cream
NuCare Pharmaceuticals, Inc.

EZRI CARE™ Clotrimazole Cream USP 1%

Drug Facts

Active ingredient

Clotrimazole 1%

Purpose

Antifungal

Uses

- For the treatment of most athlete's foot (tinea pedis), jock itch (tinea cruris), ringworm (tinea corporis)
- Relieves itching, scaling, cracking, burning, redness, soreness, irritation, discomfort and chafing associated with jock itch or itching, burning feet

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 when used for the treatment of jock itch • there is no improvement within 4 weeks when used for athlete's foot or ringworm

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

Directions

- Clean the affected area and dry thoroughly. Apply a thin layer of cream over affected area twice daily (morning and night) or as directed by a doctor • 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, consult a doctor. This product is not effective on the scalp or nails

Other information

• Do not use if seal on the tube is damaged or is not visible. To open, unscrew cap, pull tab to remove foil seal • Store at room temperature • Preserve in tight containers • See crimp of tube or carton for Lot Number and Expiry Date

Inactive ingredients

Cetostearyl Alcohol, Macrogol Cetostearyl Ether, Methylparaben, Mineral Oil, Propylene Glycol, Propylparaben, Purified Water, White Petrolatum

Questions or comments?

718-502-6610 between 9 am to 4 pm EST, Monday-Friday.

Cures Most Athlete's Foot

Compared to the active ingredient in Lotrimin[®] AF*

Relieves Itching and Burning Sensation. Relieves from Cracking and Scaling. Greaseless and non-staining.

*This product is not affiliated with, manufactured by, or produced by the makers or owners of Lotrimin[®] AF

Distributed by:

EzriCare, LLC

Lakewood, NJ

www.EzriCare.com

Made in India

Packaging

NuCare Pharmaceuticals, Inc.

NDC: 68071-3416-3
Clotrimazole 1%
30g Cream
See manufacturer's label
for full list of ingredients.

Clotrimazole 1%
Lot: 00000 NDC: 68071-3416-03
MFR NDC: 79503-0105-30 Exp.: 00-00
Serial# 0000000002

Clotrimazole 1%
Lot: 00000 NDC: 68071-3416-03
MFR NDC: 79503-0105-30 Exp.: 00-00
Serial# 0000000002

GTIN 00368071341638
Serial# 0000000002
Exp. Date 00-00
LOT#: 00000

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

Product #: R0255030

WARNING: KEEP OUT OF REACH OF CHILDREN STORE AT CONTROLLED TEMPERATURE 68-77°F.

Distributed by: EzriCare, LLC Lakewood, NJ

Packaged By: NuCare Pharmaceuticals, Inc. Orange, CA 92867

Apply every ___ hours ___ times a day.

Rev 01/01/15

CLOTRIMAZOLE CREAM 1%

clotrimazole cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-3416(NDC:79503-105)
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
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-3416-3	1 in 1 CARTON	05/25/2023	07/01/2025
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
OTC Monograph Drug	M005	08/19/2021	07/01/2025

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-3416)

