

**CLOTRIMAZOLE- clotrimazole cream
Preferred Pharmaceuticals Inc.**

**Clotrimazole Cream USP,1%
Antifungal Cream**

Drug Facts

Active ingredient

Clotrimazole 1%

Purpose

Antifungal

Uses

- cures most athlete's foot (tinea pedis), jock itch (tinea cruris), and ringworm (tinea corporis)
- for effective relief of
- itching
- scaling
- cracking
- burning
- redness
- soreness
- irritation
- discomfort

Warnings

For external use only

Ask a doctor before use

- on children under 2 years of age

When using this product

- avoid contact with eyes

Stop use and ask a doctor if

- irritation occurs
- there is no improvement within 4 weeks (for athlete's foot and ringworm) or within 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

This product is not effective on the scalp or nails.

For athlete's foot and ringworm: use daily for 4 weeks. **For jock itch:** use daily for 2 weeks.

- clean the affected area and dry thoroughly
- apply a thin layer of the product 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; change shoes and socks at least once daily

Other information

- To open: unscrew cap, use the pointed end of cap to puncture seal.
- store between 2°- 30°C (36° - 86°F)
- see carton or tube crimp for lot number and expiration date

Inactive ingredients

benzyl alcohol (1%), cetostearyl alcohol, cetyl esters wax, 2-octyldodecanol, polysorbate 60, purified water, sorbitan monostearate

Questions?

Call 1-866-923-4914

Distributed by:

Taro Pharmaceuticals

U.S.A., Inc.

Hawthorne, NY 10532

Relabeled By: Preferred Pharmaceuticals Inc.

PRINCIPAL DISPLAY PANEL - 30 g Carton

Cures Most Athlete's Foot

Athlete's Foot

Cream

Clotrimazole Cream USP,1%

Antifungal Cream

Antifungal Cream

NDC 68788-8216-3

Relabeled By: Preferred Pharmaceuticals Inc.

**Clotrimazole
Cream USP, 1%**

Generic for Lotrimin AF

Each gram contains: Clotrimazole 1%

Pkg Size: Exp Date: #####

Lot#: Batch#:

Ins:

Mfg: Taro Pharm.; Hawthorne, NY

Prod#:

Warning

Do not use on children under two years of age unless directed by a doctor. Avoid contact with the eyes. Keep out of the reach of children. For more information, see the Drug Facts, Ingredients, Warnings, Directions, and Other Information, see back of package. Store between 2°-30° C (36°-86°F).



Directions English
Apply externally _____ tim
es a day.

CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed.



GTIN

SN #####
EXP #####

Instrucciones Espanol:
Aplique
externamente _____ vece
s al dia.

Clotrimazole Cream USP, 1%
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Clotrimazole Cream USP, 1%
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Clotrimazole Cream USP, 1%
Qty:
Insurance NDC:
Lot: Bat:

Clotrimazole Cream USP, 1%
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Log

Chart

Billing

Patient

CLOTRIMAZOLE

clotrimazole cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDc:68788-8216(NDc:51672-2002)
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)	

Product Characteristics

Color	WHITE (Smooth)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8216-3	1 in 1 CARTON	09/08/2022	
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	333C	09/08/2022	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	RELABEL(68788-8216)

Revised: 1/2026

Preferred Pharmaceuticals Inc.