

TERBINAFINE HYDROCHLORIDE- terbinafine hydrochloride cream
Preferred Pharmaceuticals, Inc

Terbinafine Hydrochloride
Cream 1%
Antifungal Cream

Drug Facts

Active ingredient

Terbinafine hydrochloride 1%

Purpose

Antifungal

Uses

- cures most athlete's foot (tinea pedis)
- cures most jock itch (tinea cruris) and ringworm (tinea corporis)
- relieves itching, burning, cracking and scaling which accompany these conditions

Warnings

For external use only

Do not use

- on nails or scalp
- in or near the mouth or the eyes
- for vaginal yeast infections

When using this product do not get into the eyes. If eye contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if

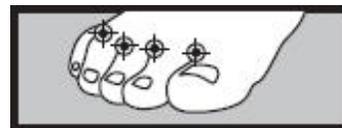
- too much irritation occurs or gets worse.
- side effects occur. You may report side effects to FDA at 1-800-FDA-1088.

Keep out of reach of children. If swallowed, get medical help or contact a poison control center right away.

Directions

- adults and children 12 years and older
 - o use the tip of the cap to break the seal and open the tube
 - o wash the affected skin with soap and water and dry completely before applying
 - o **for athlete's foot** wear well-fitting, ventilated shoes. Change shoes and socks at least once daily.

1 week between the toes



2 weeks on the bottom or sides of the foot



- **between the toes only:** apply twice a day (morning and night) for **1 week** or as directed by a doctor.
- **on the bottom or sides of the foot:** apply twice a day (morning and night) for **2 weeks** or as directed by a doctor.

- o **for jock itch and ringworm:** apply once a day (morning **or** night) for **1 week** or as directed by a doctor.
- o wash hands after each use

- children under 12 years: ask a doctor

Other information

- do not use if seal on tube is broken or is not visible
- store at controlled room temperature 20°-25°C (68°-77°F)
- see carton or tube crimp for lot number and expiration date

Inactive ingredients

benzyl alcohol, cetyl alcohol, cetyl palmitate, isopropyl myristate, polysorbate 60, purified water, sodium hydroxide, sorbitan monostearate, stearyl alcohol.

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

Terbinafine Hydrochloride

Cream 1%

Antifungal Cream

Full Prescription Strength

NET WT 1 oz (30 g)

Terbinafine Hydrochloride Cream 1%



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

Terbinafine Hydrochloride Cream 1 %
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Terbinafine Hydrochloride Cream 1 %
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Terbinafine Hydrochloride Cream 1 %
Qty: Insurance NDC:
Lot: Bat:

Terbinafine Hydrochloride Cream 1 %
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Log

Chart

Billing

Patient

Generic for Lamisil Cream
Active Ingredient Terbinafine hydrochloride 1 %

Pkg Size: Exp Date: #####/####/
Lot#: Batch#: Ins:
Mfg: Taro Pharm.; Hawthorne, NY
Prod#:

Warning
For external use only. Do not use on nails or scalp, in or near the mouth or the eyes, for vaginal yeast infections. When using this product do not get into the eyes. If eye contact occurs, rinse thoroughly with water. Stop use and ask a doctor if too much irritation occurs or gets worse, or if side effects occur. See box for additional drug facts. Keep this and all medication out of the reach of children. If swallowed, get medical help or contact a poison control center right away. Store at 20° to 25°C (68° to 77°F). Do not use if seal on tube is broken or is not visible.



Directions English
Apply externally _____ tim es a day.
Use as directed by your doctor



GTIN #####

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

Instrucciones Espanol:
Aplique externamente _____ vece s al dia.
Uso según lo dirigido por su doctor

TERBINAFINE HYDROCHLORIDE

terbinafine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-9814(NDC:51672-2080)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Terbinafine Hydrochloride (UNII: 012C11ZU6G) (Terbinafine - UNII:G7RIW8S0XP)	Terbinafine Hydrochloride	1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
benzyl alcohol (UNII: LKG8494WBH)	
cetyl alcohol (UNII: 936JST6JCN)	
cetyl palmitate (UNII: 5ZA2S6B08X)	
isopropyl myristate (UNII: 0RE8K4LNJS)	
polysorbate 60 (UNII: CAL22UVI4M)	
water (UNII: 059QF0KO0R)	
sodium hydroxide (UNII: 55X04QC32I)	
sorbitan monostearate (UNII: NVZ4I0H58X)	
stearyl alcohol (UNII: 2KR89I4H1Y)	

Product Characteristics

Color	WHITE	Score	
Shape		Size	

Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-9814-1	15 g in 1 TUBE; Type 0: Not a Combination Product	04/05/2012	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA077511	04/05/2012		

Labeler - Preferred Pharmaceuticals, Inc (791119022)

Registrant - Preferred Pharmaceuticals, Inc (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals, Inc		791119022	RELABEL(68788-9814)

Revised: 9/2025

Preferred Pharmaceuticals, Inc