

GOODSENSE TERBINAFINE HYDROCHLORIDE- terbinafine hydrochloride cream
Geiss, Destin & Dunn, Inc.

GoodSense®
Terbinafine Hydrochloride

Drug Facts

Active ingredient

Terbinafine hydrochloride 1%

Purpose

Antifungal

Uses

- cures most athlete's foot (tinea pedis)
- cures most jock itch (tinea cruris) and ring worm (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.

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
 - use the tip of the cap to break the seal and open the tube
 - wash the affected skin with soap and water and dry completely before applying
 - **for athlete's foot** wear well-fitting, ventilated shoes. Change shoes and socks at least once daily.
 - **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.
 - **for jock itch and ringworm:** apply once a day (morning **or** night) for **1 week** or as directed by a doctor.
 - 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.

Questions?

Call **1-866-923-4914**

Distributed By
Geiss, Destin and Dunn, Inc.
Peachtree City, GA 30269

PRINCIPAL DISPLAY PANEL - 15 g Tube Carton

GOODSENSE®

NDC 50804-080-01

Full Prescription Strength

Terbinafine Hydrochloride Cream 1%

Antifungal Cream

***Compare to active ingredient in
Lamisil AT®****

100%

**SATISFACTION
GUARANTEED**

NET WT 0.5 oz (15 g)

GOODSENSE®

Full Prescription Strength

Terbinafine Hydrochloride Cream 1%

Antifungal Cream

Full Prescription Strength

Terbinafine Hydrochloride Cream 1%

Antifungal Cream

• Relieves Itching & Burning • Cures Most Athlete's Foot

LPK-7538-1
1114-1
MSO

GOODSENSE®

NDC 50804-080-01

Full Prescription Strength

Terbinafine Hydrochloride Cream 1%

Antifungal Cream

NET WT 0.5 oz (15 g)

Compare to active ingredient in
Lamisil AT™

100%
SATISFACTION
GUARANTEE

NO COPY ON THIS
FLAP FOR LOT #
AND EXPIRY
DATE PRINT

Drug Facts (continued)

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2 weeks or more on the bottom or sides of the foot

1 week between the toes

• on the bottom or sides of the foot: apply twice a day (morning and night) for 2 weeks or as directed by a doctor.

• between the toes only: apply twice a day (morning and night) for 1 week or as directed by a doctor.

• for athlete's foot wear well-fitting, ventilated shoes. Change shoes and socks

• use the tip of the cap to break the seal and open the tube

• wash the affected skin with soap and water and dry completely before applying

• adults and children 12 years and older

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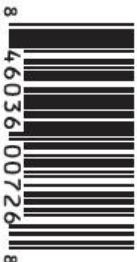
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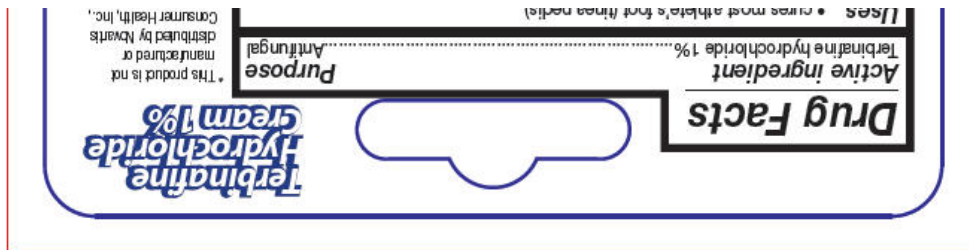


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www.lamisil.com
Peachtree City, GA 30269
Gees, Desin & Dum, Inc.
Distributed By

owner of the registered trademark Lamisil AT™

NO VARNISH
ON THIS FLAP

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ON THIS FLAP



GOODSENSE TERBINAFINE HYDROCHLORIDE

terbinafine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50804-080
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: 059QF0K00R)	
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:50804-080-01	1 in 1 CARTON		
1		15 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
ANDA	ANDA077511	07/02/2007	

Labeler - Geiss, Destin & Dunn, Inc. (076059836)

Revised: 4/2015

Geiss, Destin & Dunn, Inc.