CONAZOL ANTIFUNGAL- tolnaftate solution/ drops
MarcusUSA

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Conazol®
antifungal

Drug Facts

Active Ingredients

Tolnaftate 1%

Purposes

Antifungal

Uses

For effective treatment of athlete's foot, jock itch and ringworm. Not for relief of infected finger nails and toe nails or effective against bacteria or viruses.

Warnings

For external use only

- Avoid contact with eyes.
- **Do not use** on children under 2 years of age unless directed by a doctor.

Stop and ask a doctor if

- If treating athlete's foot and ringworm: If irritation occurs or if there is no improvement within 4 weeks.
- If treating jock itch: If irritation occurs or if there is no improvement within 2 weeks.

Keep out of the reach of children

- If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Clean or wash the affected area and dry thoroughly. Apply a thin layer of the product over the 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 toes; wear well-fitting, ventilated shoes, and change socks at least once daily. For athlete's foot and ringworm use daily for 4 weeks. If condition persists longer, consult a doctor. This product is not effective on the scalp or nails.

Other Information

- Store at room temperature 15°-30°C (59°-86°F)
- Questions or comments, call (800) 428-9489

Inactive Ingredients

Purified water, PEG-8, Cetearyl alcohol and Ceteth-20 phosphate and decetyl phosphate, Propylene glycol, Cocamidopropyl betaine, Octoxynol-9, Glyceryl stearate, Stearyl alcohol, Hydroxyethyl cellulose, Imidazolidinyl urea, Cetyl alcohol, Methylparaben, Propylparaben, Triethanolamine,Citric acid

Distributed by: MarcasUSA, LLC
El Segundo CA, 90245

PRINCIPAL DISPLAY PANEL - 30 ML Bottle Carton

NOTHING MORE EFFECTIVE

Conazol®
ANTIFUNGAL

TOE FUNGUS ELIMINATOR
with Tolnaftate 1%

Cures & Prevents Fungus

1 FL OZ (30 ML)

NDC: 75940-126-01

Clinically Proven

MAXIMUM STRENGTH WITHOUT A PRESCRIPTION

BEGINS WORKING ON CONTACT

NOTHING MORE EFFECTIVE

Conazol® ANTIFUNGAL

TOE FUNGUS ELIMINATOR

ELIMINADOR DE HONGOS DEL DEDO DEL PIE

with Tolnaftate 1%

Cures & Prevents Fungus

1 FL OZ (30 ML)

JUST 2 SIMPLE STEPS

Solo dos simples pasos

Cures & prevents fungus on the skin around, adjacent to and under the nails!

STEP 1/PASO 1:

Wash affected area with water and soap and dry thoroughly.

Lave el área afectada con agua y jabón y séquelo bien.

STEP 2/PASO 2:

Apply Conazol Toe Fungus Eliminator to affected area allowing it to absorb.

Aplique Conazol Toe Fungus Eliminator al área afectada permitiendo que se absorba.

NEW

NO-TOUCH APPLICATION

TO PREVENT THE SPREAD OF INFECTION

Conazol ANTIFUNGAL

LIOMONT

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MADE IN U.S.A.

Distributed by: MarcasUSA, LLC

TOE FUNGUS ELIMINATOR

NO-TOUCH APPLICATOR

No-contact application prevents spreading of fungus infection to other toes.

La aplicación sin contacto previene la propagación de la infección de hongos a otros dedos del pie.

1 FL OZ (30 ML)

CONAZOL ANTIFUNGAL

tolnaftate solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75940-126
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Tolnaftate (UNII: 06KB629TKV) (Tolnaftate - UNII:06KB629TKV)	Tolnaftate	10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
DIETHANOLAMINE CETYL PHOSPHATE (UNII: 4UG0316V9S)	
propylene glycol (UNII: 6DC9Q167V3)	
cocamidopropyl betaine (UNII: 5OCF3O11KX)	
octoxynol-9 (UNII: 7JPC6Y25QS)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
Stearyl alcohol (UNII: 2KR89I4H1Y)	
HYDROXYETHYL CELLULOSE (3000 MPA.S AT 1%) (UNII: 7Q6P4JN1QT)	
IMIDUREA (UNII: M629807ATL)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
methylparaben (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
TROLAMINE (UNII: 9O3K93S3TK)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75940-126-01	1 in 1 CARTON	10/29/2017	
1		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333C	10/29/2017	

Labeler - MarcusUSA (016139820)

Establishment

Name	Address	ID/FEI	Business Operations
Private Label Partners		046033481	LABEL(75940-126)

Revised: 1/2020 MarcusUSA