CLOTRIMAZOLE- clotrimazole solution The Podiatree Company

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.

Clotrimazole Topical Solution USP,1%

Drug Facts

Active ingredient

Clotrimazole 1%

Purpose

Antifungal

Uses

- cures most athlete's foot infections and ringworm infections
- relieve itching, burning, cracking, scaling and discomfort which can accompany these conditions

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 eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation occurs or there is no improvement within 4 weeks.

KEEP OUT OF REACH OF CHILDREN. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- wash affected area and dry thoroughly
- apply to the affected area twice daily (morning and night) or as directed by your 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 shoes and socks at least once daily
- for athlete's foot and ringworm infections, use daily for 4 weeks
- if condition persists longer, consult your doctor

Inactive ingredient

polyethylene glycol

Other information

• Store at 20° to 25°C (68° to 77°F)

Questions?

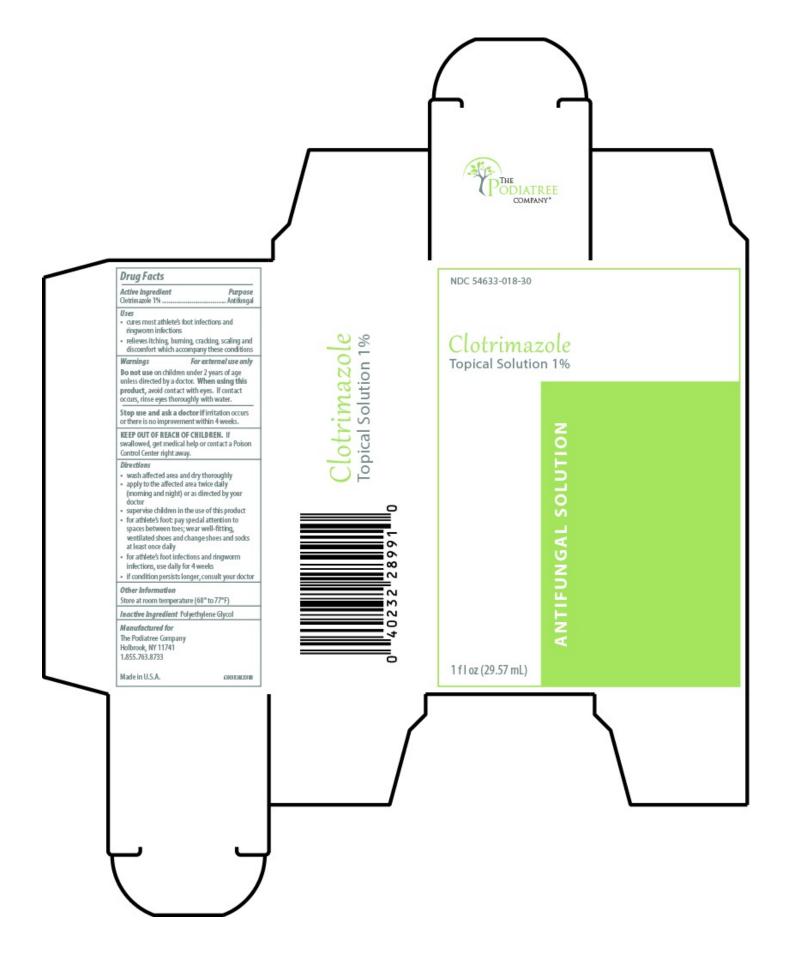
1-855-763-8733

Manufactured for

The Podiatree Company, Holbrook, NY 11741

Made in U.S.A.

PRINCIPAL DISPLAY PANEL - 30 mL Carton



clotrimazole solution

Product	Information
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:54633-018

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength		
CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)	CLOTRIMAZOLE	1g in 1mL		

Inactive Ingredients

POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)

Packaging#Item CodePackage DescriptionMarketing Start DateMarketing End Date1NDC:54633-018-301 in 1 CARTON10/26/2018

1 30 mL in 1 BOTTLE; 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	10/26/2018	

Labeler - The Podiatree Company (078656000)

Revised: 10/2018 The Podiatree Company