CLOTRIMAZOLE ANTIFUNGAL- clotrimazole solution Trifecta Pharmaceuticals USA, LLC.

Globe Clotrimazole (Topical Solution) USP 1%

Active Ingredient

Clotrimazole USP, 1.0%

Purpose

Antifungal

Uses

Cures most

- Athlete's foot (tinea pedis)
- Jock Itch (Tinea Cruris)
- Ringwork (Tinea corporis)

Effectively relieves

- Itching
- Cracking
- Burning

Discomfort that can accompany these conditions

Warnings

For External Use Only

Stop use and ask a doctor

- If irritation occurs
- If there is no improvement within 4 weeks (for athlete's foot and ringworm) or 2 weeks (for jock itch)

Ask a Doctor before use

Ask a doctor before use on children under 2 years of age

When using this product

Avoid Contact with Eyes.

Keep out of the reach of children

If swallowed, get medical help or contact a Poison Control Center immediately

Directions

- Spray over affected area
- If bandaged, let dry first
- Not to be used on children under 12 years of age

Other information

• Store at controlled room temperature 15°-30°C (59°-86°F)

Inactive Ingredients

Ethanol, Propylene glycol, purified water

Questions or Comments

Call 1-888-296-9067

Distributed By:

Trifecta Pharmaceuticals USA®

101 NE Third Avenue, Suite 1500

Ft. Lauderdale, FL. 33301 USA

www.trifecta-pharma.com

Globe® Clotrimazole anti-fungal solution is not manufactured or distributed by Bayer, owner of the registered trademark Lotrimin®AF.

Packaging

OUTSIDE BOX





INNER LABEL



CLOTRIMAZOLE ANTIFUNGAL

clotrimazole solution

Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69396-143			
Route of Administration	TOPICAL					

Active Ingredient/Active Moiety Ingredient Name CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65) CLOTRIMAZOLE (UNII: G07GZ97H65) CLOTRIMAZOLE 10 g in 1 L

Inactive Ingredients				
Ingredient Name	Strength			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
ALCOHOL (UNII: 3K9958V90M)				

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69396-143- 33	1 in 1 BOX	11/13/2023			
1		0.01 L 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 Drug	M005	11/13/2023			

Labeler - Trifecta Pharmaceuticals USA, LLC. (079424163)

Revised: 2/2024 Trifecta Pharmaceuticals USA, LLC.