WELMATE CLOTRIMAZOLE TOPICAL SOLUTION- clotrimazole liquid
YYBA CORP

WELMATE Clotrimazole Topical Solution

Drug Facts

Active ingredient

Clotrimazole USP, 1%

Purpose

Antifungal

Uses

Cures most
- Athlete's foot (tinea pedis)
- Jock Itch (tinea cruris)
- Ringworm (tinea corporis) Effectively relieves • Itching • Cracking • Burning Discomfort which can accompany these conditions

Warnings
- **For external use only**

Ask a doctor before use

on children under 2 years of age

When using this product

avoid contact with eyes

Stop use and ask a doctor if
- irritation occurs
- there is no improvement within 4 weeks (for athlete's foot and ringworm) or 2 weeks (for jock itch)

Keep out of reach of children

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

Directions
- This product is not effective on the scalp or nails. For best results, follow directions and continue treatment for length of time indicated. For athlete's foot and ringworm: use daily for 4 weeks. For jock itch: use daily for 2 weeks.

- clean the affected area and dry thoroughly
- apply a thin layer of the product over 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 the toes; wear well-fitting, ventilated shoes; change shoes and socks at least once daily.

Other information

Store at 15° to 30°C (59° to 86°F)

Inactive ingredient

polyethylene glycol 400

Questions?

call 866-933-6337

Package Labeling:

WELMATE CLOTRIMAZOLE TOPICAL SOLUTION

clotrimazole liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73581-449
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)	CLOTRIMAZOLE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73581-449-00	1 in 1 BOX	02/19/2024	
1		10 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 Drug	M005	02/19/2024	

Labeler - YYBA CORP (006339772)

Revised: 2/2024 YYBA CORP