CLOTRIMAZOLE- drs. clotrimazole cream OL PHARMA TECH,LLC

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.

ACTIVE INGREDIENT

Clotrimazole 1%

PURPOSE

Antifungal

USES

- cures most athlete's foot, jock itch and ringworm
- relieves itching, burning, cracking, scaling and discomfort which accompany these conditions

warnings

For external use only

Do not use:

Do not use on children under 2 years of age unless directed by a doctor.

WHEN USING THIS PRODUCT

avoid contact with the eyes

Stop use and ask your doctor if:

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

- wash 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,

and change shoes and socks at least once daily

for athlete's foot and ringworm, use daily for 4 weeks; for jock itch, use daily for 2 weeks

if condition lasts longer, contact a doctor

• this product is not effective on the scalp or nails

Other information

- store at controlled room temperature 59°-86°F (15°-30°C).
- before using any medication, read all label directions. Keep carton, it contains important information

Inactive ingredients

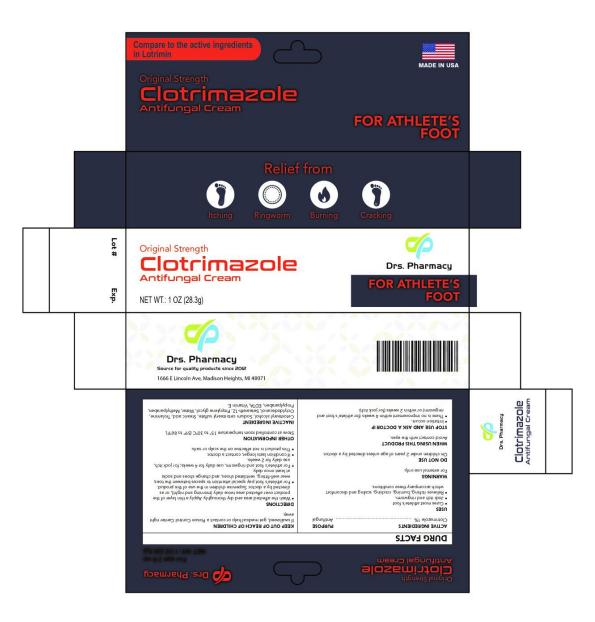
sodium cetostearyl sulfate, cetostearyl alcohol, stearic acid, trolamine, mineral oil , ceteareth-12, propylene glycol, methyl parabeen, propyl paraben, EDTA disodium, vitamin E, water

For Questions:

www.drspharmacyusa.com

original strenghth

Antifungal cream



CLOTRIMAZOLE						
drs. clotrimazole cream						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:80489-002		
Route of Administration	TOPICAL					
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Active Ingredient/Active Moiety						
Ingredient Name			Basis of Strength		Stre	ngth
CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)		CLOTRIMAZ OLE		10 mg	in 1 g	
Inactive Ingredients						

Ingredient Name	Strength
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CETEARETH-12 (UNII: 7V4MR24V5P)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
.ALPHATOCOPHEROL (UNII: H4N855PNZ1)	
MINERAL OIL (UNII: T5L8T28FGP)	
SODIUM CETOSTEARYL SULFATE (UNII: 7ZBS06BH4B)	
TROLAMINE (UNII: 903K93S3TK)	
WATER (UNII: 059QF0KO0R)	
PROPYLPARABEN (UNII: Z8IX2SC10H)	

Product Characteristics				
Color	white	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				

Packaging

	5 5				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:80489-002- 01	1 in 1 CARTON	01/01/2021		
1 28.3 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	
от	C monograph fina	I part333C	01/01/2021		

Labeler - OL PHARMA TECH,LLC (021170377)

Registrant - OL PHARMA TECH,LLC (021170377)

Establishment

Name	Address	ID/FEI	Business Operations
OL PHARMA TECH,LLC (Drs, pharmacy)		021170377	manufacture(80489-002)

Revised: 10/2023

OL PHARMA TECH,LLC