QUALITY CHOICE CLOTRIMAZOLE- clotrimazole cream
Chain Drug Marketing Association

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.

QC Clotrimazole Cream 1 oz 99261, 2019

Active ingredient Purpose

Clotrimazole 1%..... Antifungal

Uses
- cures most athlete's foot, jock itch, and ringworm

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 the eyes

Stop use and ask a doctor if
- irritation occurs
- there is no iprovement 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 immediately.

Directions
- wash affected area and dry thoroughly
- apply a thin layer over affected area twice daily (morning and night)
- supervise children in the use of this product
- for athlete's foot: pay special attention to spaces between the toesl; wear well-fitting, ventilated shoes and change shoes and socks at least once daily
- if condition persists longer, ask a doctor
- this product is not effective on the scalp or nails

Other information
- store between 20° to 25°C (68° to 77°F)
- Lot No. & Exp. Date: see box or see crimp of tube

Inactive ingredients benzyl alcohol, cetanol, liquid paraffin, octyldodecanol, polysorbate 60, propylene glycol, purified water, sodium hydroxide, sodium monostearate, stearic acid, stearyl alcohol

DISTRIBUTED BY C.D.M.A., INC.

43157 W. NINE MILE

NOVI, MI 48376-0995

www.qualitychoice.com

MADE IN KOREA

NDC 63868-595-28

*Compare to the active ingredient in LOTRIMIN®

Clotrimazole Cream

Cures Most Athlete's Foot

Antifungal Cream 1% USP

Relieves Itching, Burning, Cracking & Scaling

Clotrimazole Cream

Cures Most Athlete's Foot

Antifungal Cream 1% USP

1 oz NET WT (28g)

*This product is not manufactured or distributed by Bayer Consumer Care, Inc., owner of the registered trademark Lotrimin®.

Made in Korea

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43157 W. Nine Mile
Novi, MI 48376-0995
www.qualitychoice.com
Questions: 248-449-9300

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Clotrimazole Cream
Cures Most Athlete's Foot

LOT & EXP.

Drug Facts

Active ingredient — **Purpose**

Clotrimazole 1% ..... 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 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 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 immediately.

Directions

- wash affected area and dry thoroughly
- apply a thin layer over affected area twice daily (morning and night)
- 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 persists longer, ask a doctor
- this product is not effective on the scalp or nails

Other information ■ store between 20° to 25°C (68° to 77°F) ■ Lot No. & Exp. Date: see box or see crimp of tube

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Questions or comments? 1-800-814-8028

QUALITY CHOICE CLOTRIMAZOLE

clotrimazole cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-595
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 g

Inactive Ingredients	
Ingredient Name	**Strength**
BENZYL ALCOHOL (UNII: LKG8494WBH)	
PARAFFIN (UNII: I9O0E3H2ZE)	
OCTYLDODECANOL (UNII: 461N1O614Y)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63868-595-28	1 in 1 CARTON	01/20/2015	
1		28 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
OTC monograph final	part333C	01/19/2015	

Labeler - Chain Drug Marketing Association (011920774)

Revised: 12/2019 Chain Drug Marketing Association