# FAMILY CARE TOLNAFTATE - tolnaftate cream United Exchange Corp.

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.

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# Family Care Tolnaftate Cream 1oz (502 TG, 2018)

# Active ingredient Purpose

Tolnaftate 1%...... Antifungal

#### Uses

- proven clinically effective in the treatment of athlete's foot (tinea pedis) and ringworm (tinea corporis)
- helps prevent most athlete's foot with daily use
- for effective relief of itching, burning and cracking

# **Warnings**

For external use only

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

Do not use on children under 2 years of age except under the advice and supervision of a doctor

**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
- use daily for 4 weeks; if condition persists longer, ask a doctor
- to prevent athlete's foot, apply once or twice daily (morning and/or night)
- this product is not effective on the scalp or nails

#### Other information

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

**Inactive ingredients** ceteth-20, cetostearyl alcohol, chlorocresol, liquid paraffin, monobasic sodium, phosphate, dihydrate, propylene glycol, purified water, white petrolatum

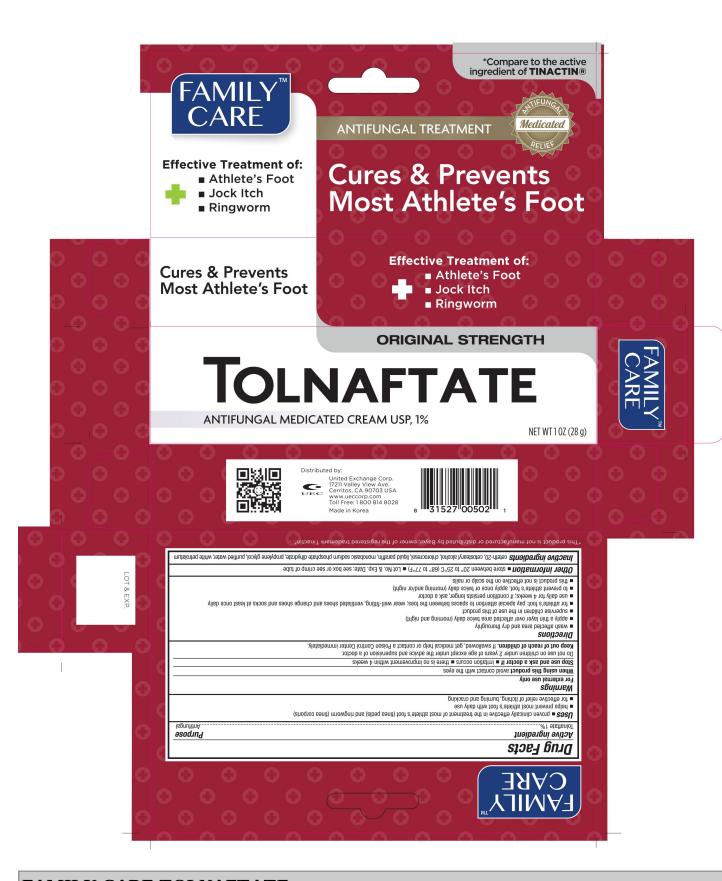
#### Distributed By:

United Exchange Corp.

17211 Valley View Ave.

Cerritos, CA 90703 USA

Made in Korea



# **FAMILY CARE TOLNAFTATE**

tolnaftate cream

<b>Product Information</b>	roduct Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-502			
Route of Administration	TOPICAL					

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	10 mg in 1 g		

Inactive Ingredients			
Ingredient Name	Strength		
CETETH-20 (UNII: 1835H2IHHX)			
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)			
CHLOROCRESOL (UNII: 36W53O7109)			
PARAFFIN (UNII: 19 O 0 E 3 H 2 Z E )			
SODIUM PHO SPHATE, MO NO BASIC, DIHYDRATE (UNII: 5QWK665956)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
PETROLATUM (UNII: 4T6H12BN9U)			

P	Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date	
1	NDC:65923-502-28	1 in 1 CARTON	10/05/2016		
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	04/30/2013			

# Labeler - United Exchange Corp. (840130579)

Revised: 12/2018 United Exchange Corp.