SUNMARK ANTIFUNGAL- tolnaftate cream
A-S Medication Solutions

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sunmark™
antifungal

Drug Facts

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- clinically proven to cure most athlete's foot (tinea pedis), jock itch (tinea cruris) and ringworm (tinea corporis)
- helps prevent most athlete's foot from recurring when used daily
- effectively soothes and relieves symptoms of athlete's foot, including 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 (for athlete's foot or ringworm) or within 2 weeks (for jock itch)

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 right away.

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
- this product is not effective on scalp or nails

For athlete's foot

- pay special attention to spaces between toes
- wear well-fitting shoes, 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, consult a doctor

Other information
- Store between 15° - 30°C (59° - 86°F)
- See end panel of carton and tube crimp for lot number and expiration date

Inactive ingredients

BHT, PEG-400, PEG-3350, titanium dioxide, white petrolatum

Distributed by McKesson
One Post Street
San Francisco, CA 94104

HOW SUPPLIED

Product: 50090-3358

NDC: 50090-3358-0 15 g in a TUBE / 1 in a CARTON

Tolnaftate

SUNMARK ANTIFUNGAL
tolnaftate cream
Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50090-3358(NDC:49348-155)
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
Butylated Hydroxytoluene (UNII: 1P9D0Z171K)	
Polyethylene Glycol 400 (UNII: B697894SGQ)	
Polyethylene Glycol 3350 (UNII: G2M7P15E5P)	
titanium dioxide (UNII: 15FIX9V2JP)	
petrolatum (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-3358-0	1 in 1 CARTON	02/01/2018	
1		15 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 Drug	M005	02/17/2006	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-3358)

Revised: 12/2023 A-S Medication Solutions