

DERMAN ANTIFUNGAL CREAM- tolnaftate cream
Taisho Pharmaceutical California Inc.

Derman antifungal cream

Drug Facts

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- cures most athlete's foot, jock itch, and ringworm
- relieves itching, burning, cracking, scaling, redness, soreness, irritation and discomfort

Warnings

Do not use

on children under 2 years of age unless directed by a doctor.

For external use only.

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

If pregnant or breast-feeding,
ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away: 800-222-1222.

Directions

- wash affected area and dry thoroughly
- apply a thin layer 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 persists longer, consult a doctor
- this product is not effective on the scalp or nails

Other information

store at 15 to 30°C (59° to 86°F)

DO NOT USE IF SEAL ON TUBE IS PUNCTURED OR MISSING. **Tamper Evident:**

Inactive ingredients

ceteth-20, cetyl alcohol, chlorocresol, mineral oil, monobasic sodium phosphate, petrolatum, propylene glycol, purified water, stearyl alcohol

Package Labeling:25g

NO
COLOR

Drug Facts (continued)

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PL33102B
V01
NO
COLOR

ANTIFUNGAL CREAM
DERMAN[®]
Tolnaftate 1%

DERMAN[®]

NDC 81929-004-02

ANTIFUNGAL CREAM
Tolnaftate 1%

Cures And Prevents Most Athletes Foot
Relieves Itching And Burning



NET WT. 0.88 OZ (25 g)

NO
COLOR



NO
COLOR

DISTRIBUTED BY

Taisho Pharmaceutical California Inc. 3528 Torrance Blvd, Suite #320 Torrance, CA 90503 U.S.A.

Drug Facts

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Tolnaftate 1%.....Antifungal

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GLUE FLAP
NO COLOR / TEXT
NO COATING

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NO COLOR / TEXT
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NDC 81929-004-02

DERMAN[®]

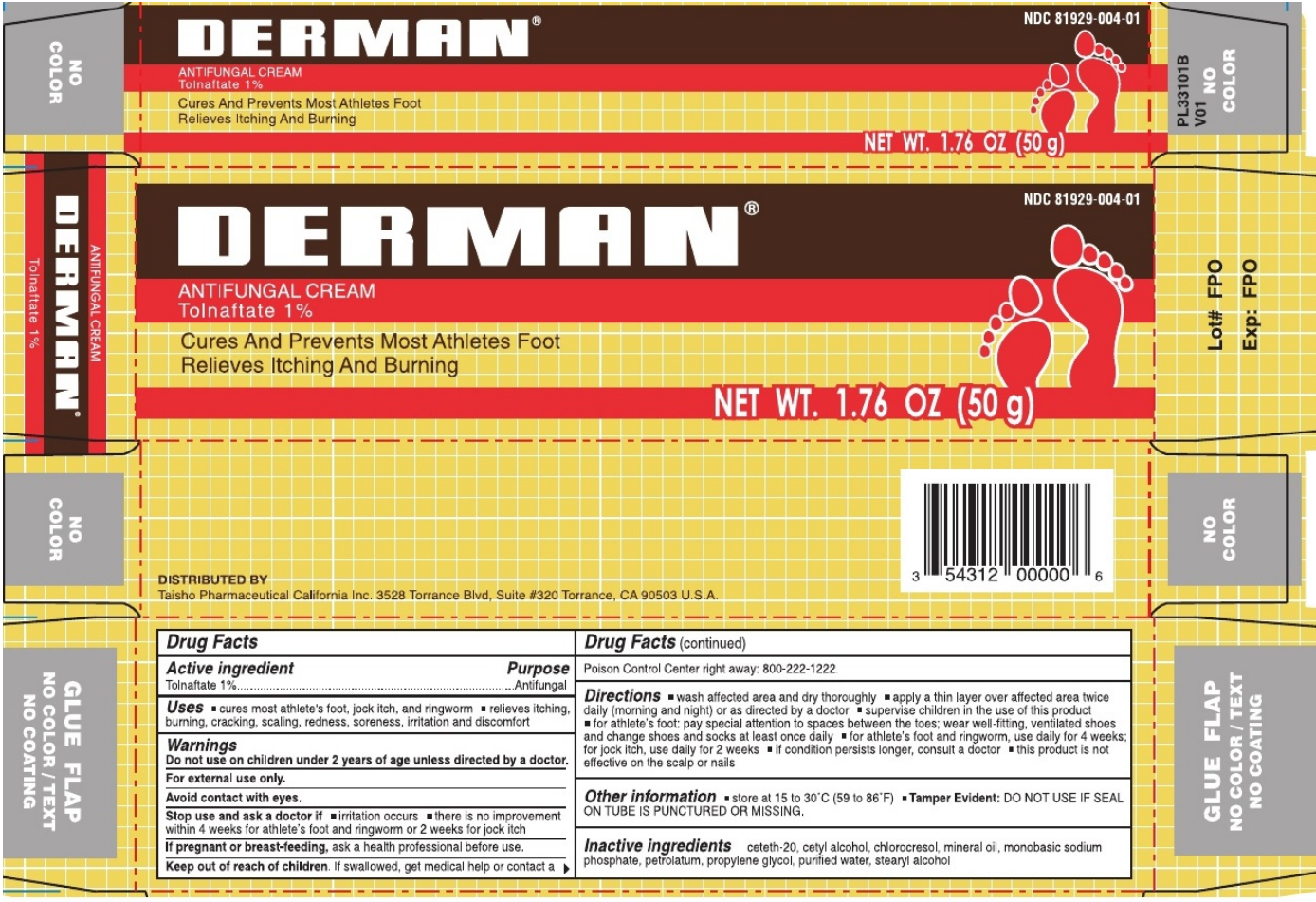
ANTIFUNGAL CREAM
Tolnaftate 1%

FOR THE TREATMENT OF ATHLETE'S FOOT
NET WT. 0.88 OZ (25 g)



Drug Facts		Drug Facts (continued)	
Active ingredient Tolnaftate 1%.....	Purpose Antifungal	right away: 800-222-1222.	
Uses ■ cures most athlete's foot, jock itch, and ringworm ■ relieves itching, burning, cracking, scaling, redness, soreness, irritation and discomfort		Directions ■ wash affected area and dry thoroughly ■ apply a thin layer 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 persists longer, consult a doctor ■ this product is not effective on the scalp or nails	
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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		PL33102T V01	
If pregnant or breast-feeding, ask a health professional before use.			
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center ▶			

Package Labeling:50g



DERMAN[®]

NDC 81929-004-01

ANTIFUNGAL CREAM
Tolnaftate 1%

Cures And Prevents Most Athletes Foot
Relieves Itching And Burning

NET WT. 1.76 OZ (50 g)



PL33101B
V01 NO
COLOR

DERMAN[®]

NDC 81929-004-01

ANTIFUNGAL CREAM
Tolnaftate 1%

Cures And Prevents Most Athletes Foot
Relieves Itching And Burning

NET WT. 1.76 OZ (50 g)



Lot# FPO
Exp: FPO



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Tolnaftate 1%

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ANTIFUNGAL CREAM
Tolnaftate 1%

FOR THE TREATMENT OF ATHLETE'S FOOT
NET WT. 1.76 OZ (50 g)



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		NDC 31929-004-01 PL33101T V01

DERMAN ANTIFUNGAL CREAM

tolnaftate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81929-004
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: I835H2IHHX)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CHLOROCRESOL (UNII: 36W53O7109)	
MINERAL OIL (UNII: T5L8T28FGP)	

SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JH2SW)	
PETROLATUM (UNII: 4T6H12BN9U)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81929-004-02	1 in 1 CARTON	07/01/2022	
1		25 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:81929-004-01	1 in 1 CARTON	07/01/2022	
2		50 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	07/01/2022	

Labeler - Taisho Pharmaceutical California Inc. (603827635)

Revised: 11/2023

Taisho Pharmaceutical California Inc.