DERMAN ANTIFUNGAL- tolnaftate aerosol
Taisho Pharmaceutical California Inc.

Derman Antifungal Liquid Spray

Drug Facts

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Uses

- Clinically proven to prevent most athlete’s foot.
- prevent the recurrence of athlete's foot with daily use
- for effective relief of itching, cracking and burning

Warnings

For external use only

Flammable: Contents under pressure. Avoid spraying in eyes. Do not use or store near heat or open flame. Do not puncture or incinerate container. Do not store at temperature above 120ºF. Keep out of the reach of children. Use only as directed. Intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal.

When using this product

- avoid contact with the eyes or mouth
- use only as directed

Stop use and ask a doctor if

- irritation occurs
- no improvement within four weeks for athlete's foot and ringworm

Do not use

on children under 2 years of age unless directed by 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
- shake can well and spray 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
- if condition persists, consult a doctor
- to prevent athlete's foot: wash the feet and dry thoroughly; spray a thin layer of the product to the feet once or twice daily (morning and/or night)
- this product is not effective on the scalp or nails
- if nozzle clogs, clean with a pin

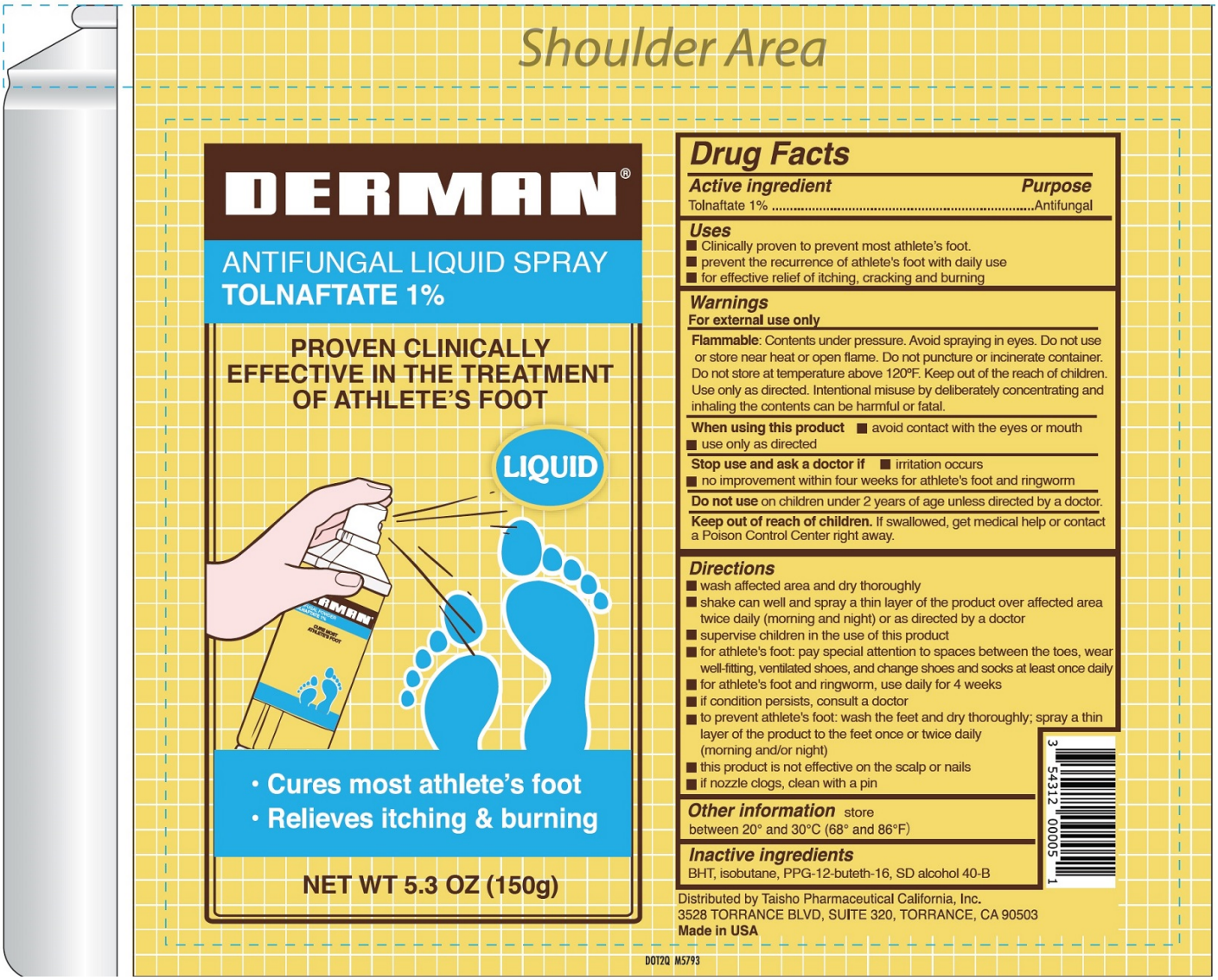
Other information

store between 20° and 30°C (68° and 86°F)

Inactive ingredients

BHT, isobutane, PPG-12-buteth-16, SD alcohol 40-B

Package Labeling:



DERMAN ANTIFUNGAL

tolnaftate aerosol

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81929-006
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)	
ISOBUTANE (UNII: BXR49TP611)	
PPG-12-BUTETH-16 (UNII: 58CG7042J1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81929-006-01	150 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	10/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	10/01/2024	

Labeler - Taisho Pharmaceutical California Inc. (603827635)

Revised: 5/2024 Taisho Pharmaceutical California Inc.