LOTRIMIN DAILY PREVENT DEODORANT- tolnaftate aerosol, powder Bayer HealthCare LLC.

Lotrimin Daily Prevention Powder Spray TALC FREE UI 1615422

Drug Facts

Active Ingredient

(To Deliver)Tolnaftate 1%

Purpose

Antifungal

Use

Use

clinically proven to prevent most athlete's foot with daily use

Warnings

For external use only

Flammable: Do not use near heat, flame, or while smoking

Do not use on children under 2 years of age unless directed by a doctor

When using this product

- avoid contact with the eyes
- use only as directed. Intentional misuse by deliberately concentrating and inhaling contents can be harmful or fatal.
- contents under pressure. Do not puncture or incinerate. Do not store at temperatures above 120°F.

Stop use and ask a doctor if irritation occurs.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Directions

- to prevent athlete's foot, wash the feet and dry thoroughly.
- shake can well and spray a thin layer of the product on the feet once or twice daily (morning and/or night).
- supervise children in the use of this product.
- pay special attention to spaces between the toes; wear well-fitting, ventilated shoes, and change shoes and socks at least once daily.

Other information

Other Information

store between 20° to 25°C (68°- 77°F)

Inactive Ingredients

butylated hydroxytoluene, fragrance, hydroxypropyl cellulose, isobutane, kaolin, magnesium stearate, PPG-12-Buteth-16, SD alcohol 40B (9% w/w), zea mays (corn) starch

Questions?1-866-360-3266 or visit us at www.lotrimin.com

Display can 160 grams



LOTRIMIN DAILY PREVENT DEODORANT

tolnaftate aerosol, powder

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11523-0136
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	
MAGNESIUM STEARATE (UNII: 70097M6I30)		
STARCH, CORN (UNII: O8232NY3SJ)		
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)		
ALCOHOL (UNII: 3K9958V90M)		
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)		
PPG-12-BUTETH-16 (UNII: 58CG7042J1)		
ISOBUTANE (UNII: BXR49TP611)		
KAOLIN (UNII: 24H4NWX5CO)		

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:11523- 0136-1	160 g in 1 CAN; Type 0: Not a Combination Product	03/01/2024		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	03/01/2024	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 12/2025 Bayer HealthCare LLC.