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 120ºF.

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 20º to 25ºC (68º– 77ºF)

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: 6/2024 Bayer HealthCare LLC.