LOTRIMIN DAILY PREVENTION DEO- tolnaftate aerosol, powder Bayer HealthCare LLC.

Lotrimin Daily Prevention Deo Powder Spray UI 1612507

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

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.

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.

do not puncture or incinerate. Contents under pressure. Do not store at temperatures above 120°F

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

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store between 20° to 25° C (68° – 77° F)

Inactive Ingredients

butylated hydroxytoluene, fragrance, isobutane, PEG-12-Buteth-16, SD alcohol 40B (10.5% v/v), talc

Questions? 1-866-360-3266

Display can 160 grams

Lotrimin ® AF

TOLNAFTATE ANTIFUNGAL

deodorant powder spray

DAILY PREVENTION

clinically proven to

prevent most

athlete's foot

- stops the growth of most athlete's foot fungus
- instantly cools and dries feet
- fights odor all day

NET WT 160 g (5.6 OZ)

LOT CODE AREA NO PRINT IN THIS AREA BACK PANEL ONLY





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Dist. by: Bayer HealthCare LLC Whippany, NJ 07981 87045560

Bayer

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2D Code

LOTRIMIN DAILY PREVENTION DEO

tolnaftate aerosol, powder

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11523-0010

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
ALCOHOL (UNII: 3K9958V90M)	
TALC (UNII: 7SEV7J4R1U)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PPG-12-BUTETH-16 (UNII: 58CG7042J1)	
ISOBUTANE (UNII: BXR49TP611)	

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:11523- 0010-1	160 g in 1 CAN; Type 0: Not a Combination Product	02/14/2020				
2	NDC:11523- 0010-2	133 g in 1 CAN; Type 0: Not a Combination Product	02/14/2020				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M005	02/14/2020		

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 12/2025 Bayer HealthCare LLC.