LOTRIMIN DAILY PREVENTION- tolnaftate powder
Bayer HealthCare LLC.

Lotrimin Daily Prevention Powder UI 1612510

Drug Facts

Active ingredient

Tolnaftate 1%

Purpose

Antifungal

Use

Use

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

Warnings

For external use only

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

When using this product avoid contact with the eyes

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.

• apply 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 socks at least

once daily.

Other information

Other information

store between 20º to 25ºC (68º– 77ºF)

Inactive ingredients

benzethonium chloride;corn starch, kaolin;sodium bicarbonate

Questions

Questions? 1-866-360-3266

Package Display 3 oz. label

LOTRIMIN® AF

TOLNAFTATE ANTIFUNGAL

medicated foot powder

DAILY

PREVENTION

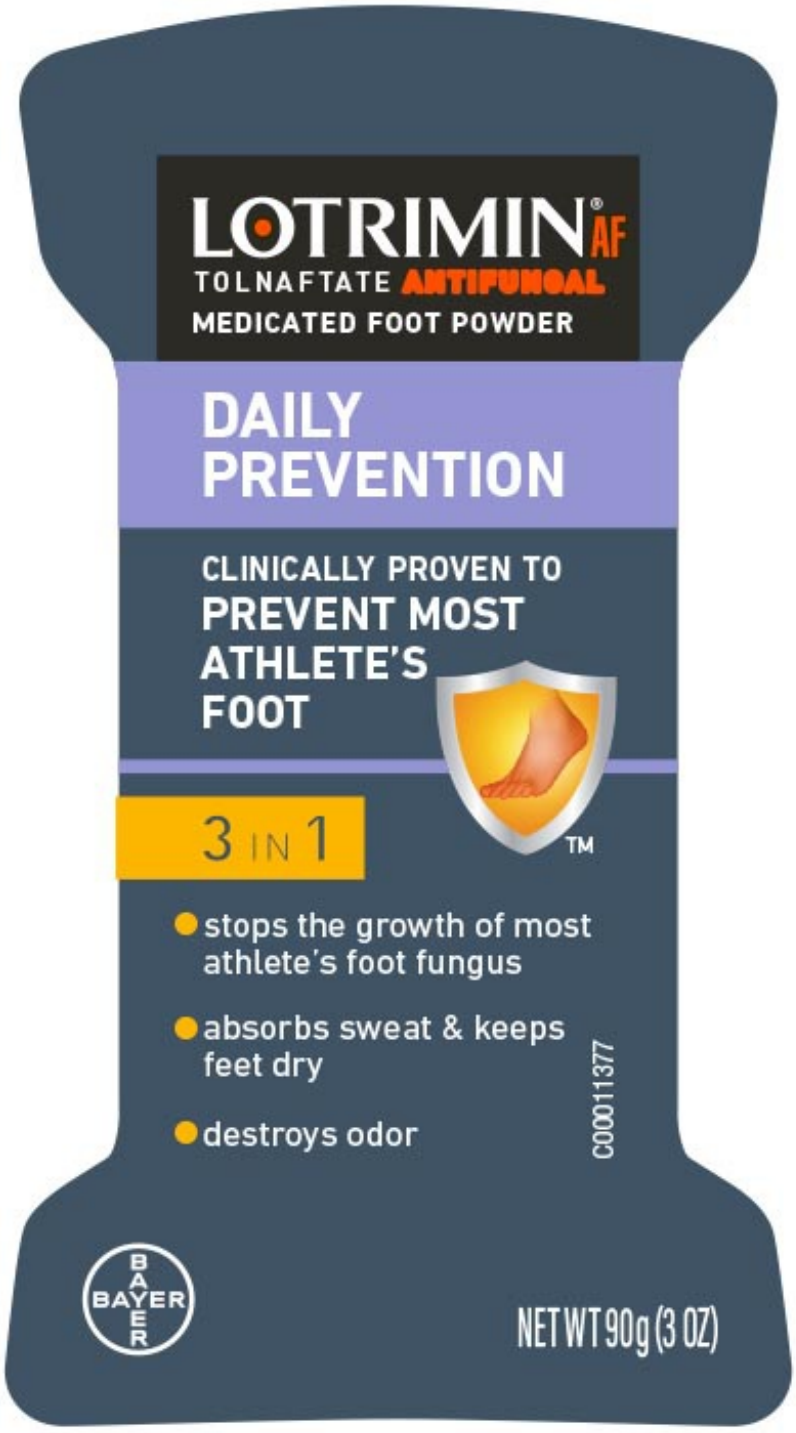
clinically proven to

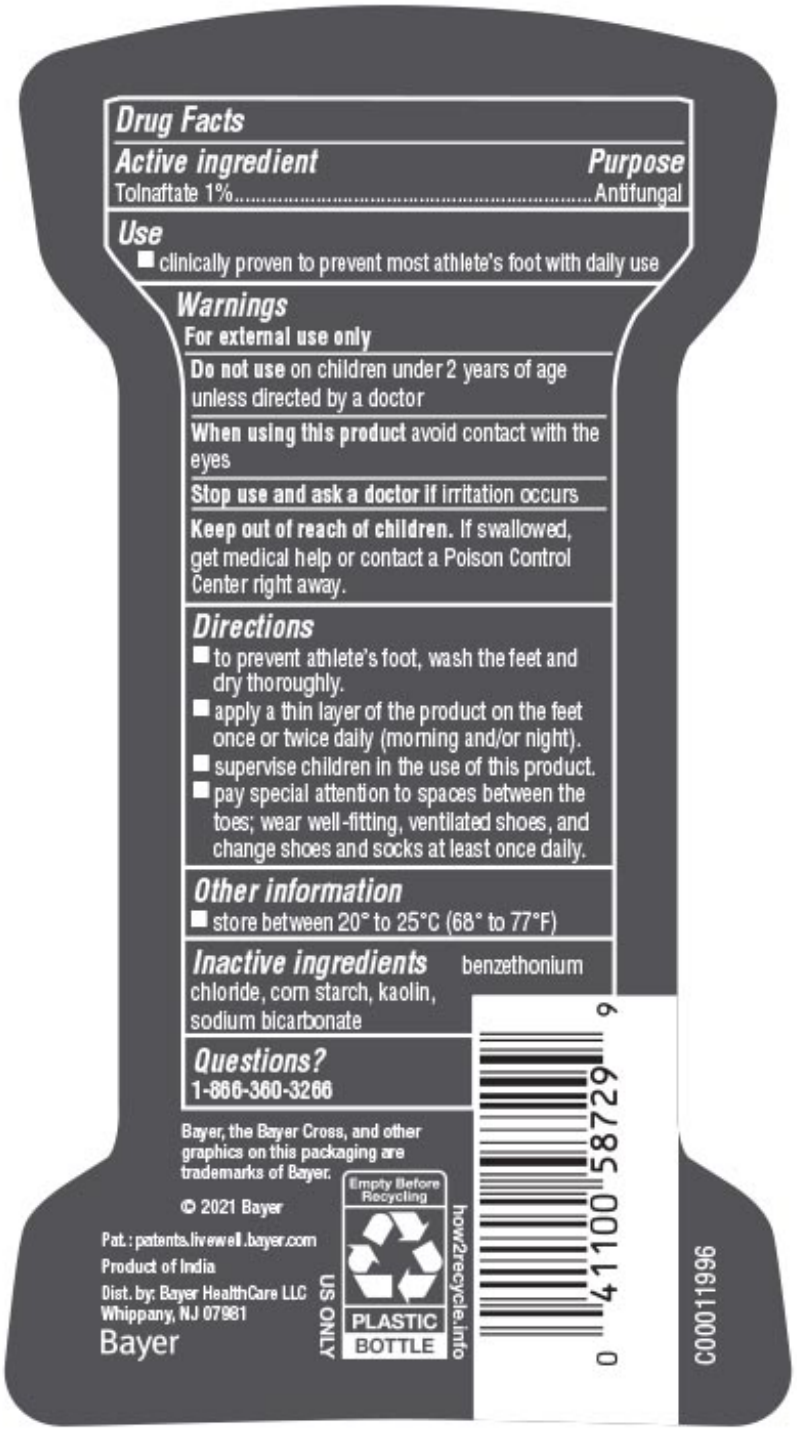
prevent most

athlete's foot

- stops the growth of most athlete's foot fungus
- absorbs sweat & keeps feet dry
- destroys odor

NET WT 90g (3 OZ)


LOTRIMIN® AF
TOLNAFTATE ANTIFUNGAL
MEDICATED FOOT POWDER
DAILY PREVENTION
CLINICALLY PROVEN TO PREVENT MOST ATHLETE'S FOOT
3 IN 1
™
stops the growth of most athlete's foot fungus
absorbs sweat & keeps feet dry
destroys odor
C00011377
BAYER
NET WT 90g (3 OZ)



LOTRIMIN DAILY PREVENTION

tolnaftate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-0011
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
KAOLIN (UNII: 24H4NWX5CO)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
STARCH, CORN (UNII: O8232NY3SJ)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-0011-1	90 g in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	
2	NDC:11523-0011-2	28 g in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2022	

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: 12/2024 Bayer HealthCare LLC.