LOTRIMIN DAILY PREVENTION- tolnaftate powder
Bayer HealthCare LLC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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

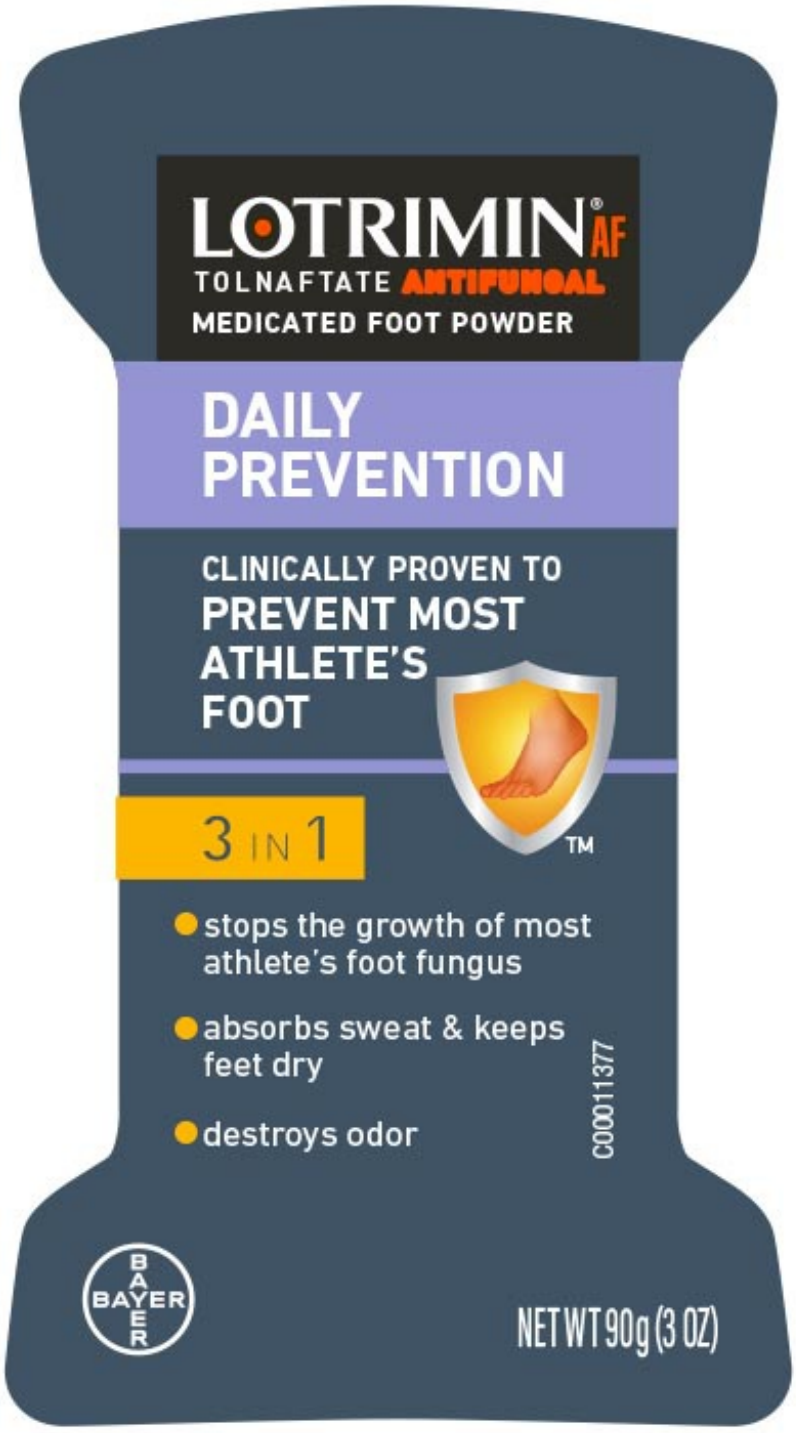
clinically proven to

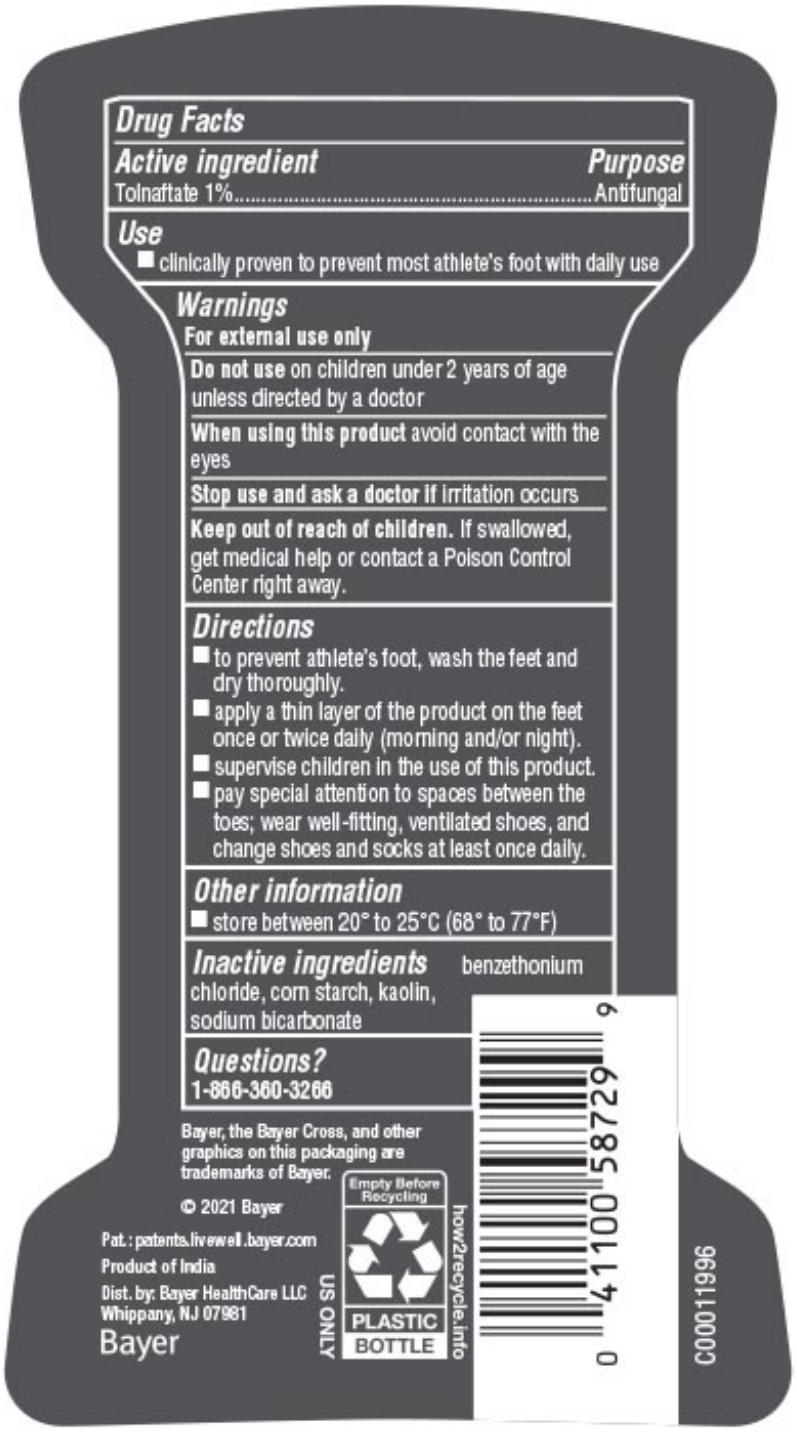
prevent most

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 final	part333C	02/14/2020	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 9/2023 Bayer HealthCare LLC.