LOTRIMIN DAILY SWEAT AND ODOR CONTROL- topical starch powder
Bayer HealthCare LLC.

Lotrimin Daily Sweat and Odor Control Powder UI 1612345

Drug Facts

Active ingredient

Topical starch 83.7%

Purpose

Skin Protectant

Use

Use

temporarily protects and helps relieve minor skin irritation

Warnings

For external use only

Do not use on broken skin.

When using this product

• Do not get into eyes

• Keep away from face and mouth to avoid breathing it

Stop use and ask a doctor if

- Condition worsens
- Symptoms last more than 7 days or clear up and occur again within a few days

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

Directions

Directions

Apply as needed

Inactive ingredients

benzethonium chloride,camphor,eucalyptus globulus leaf oil, fragrance, kaolin,lemon oil, zinc oxide

Questions

Questions? 1-866-360-3266

Package display

LOTRIMIN

TOPICAL STARCH SKIN PROTECTANT

medicated foot powder

DAILY SWEAT

& ODOR CONTROL

6 odor-fighting

ingredients to control odor

- super sweat absorbent powder
- relieves irritation
- talc-free

NET WT 177 G (6.25 OZ)

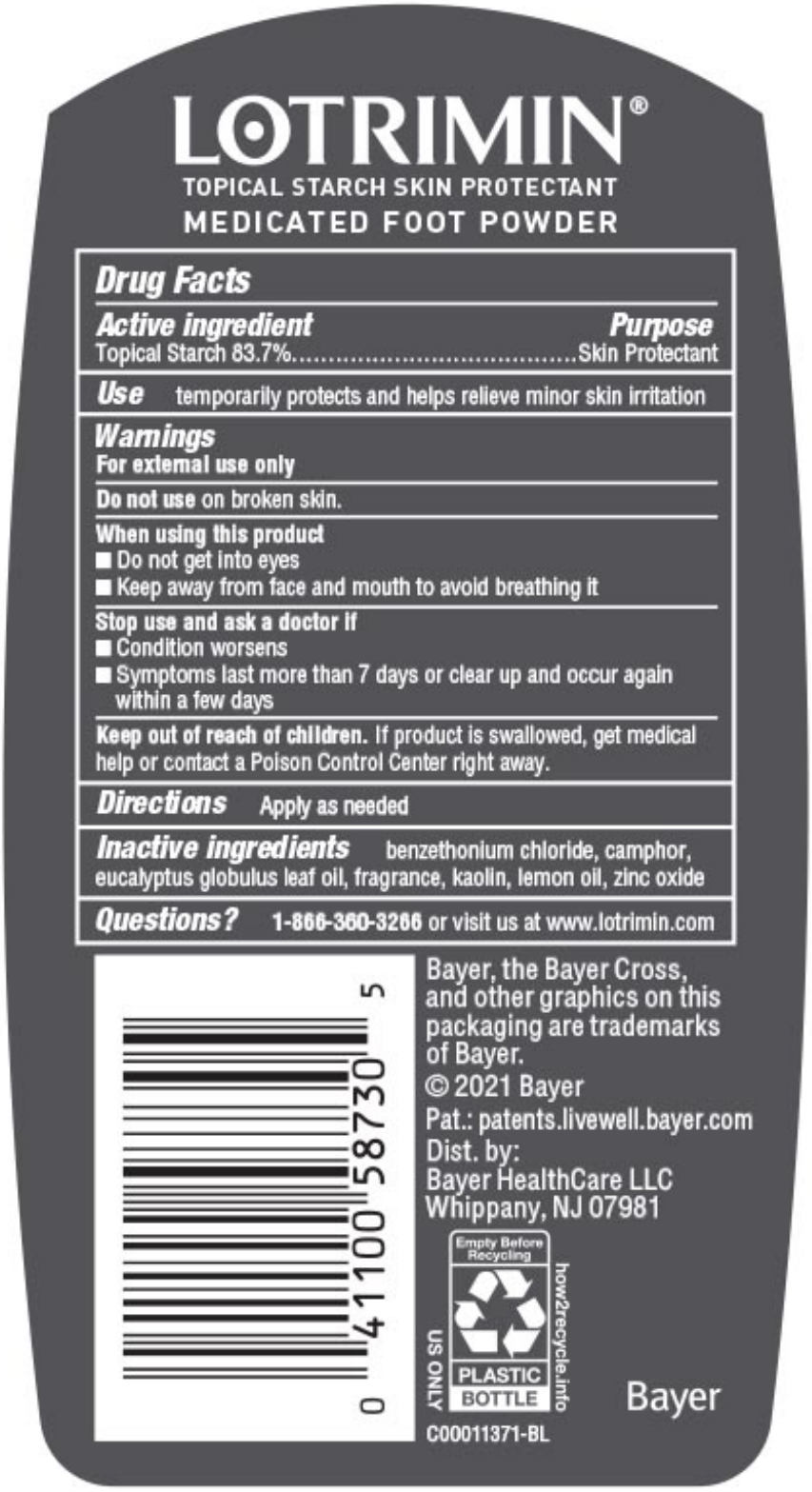
Relieves irritation

Talc-free

C00011371-FL

BAYER

NET WT 177g (6.25 OZ)



LOTRIMIN DAILY SWEAT AND ODOR CONTROL

topical starch powder

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-0012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
STARCH, CORN (UNII: O8232NY3SJ) (STARCH, CORN - UNII:O8232NY3SJ)	STARCH, CORN	837 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
EUCALYPTUS OIL (UNII: 2R04ONI662)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	
ZINC OXIDE (UNII: SOI2LOH54Z)	
LEMON OIL (UNII: I9GRO824LL)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
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:11523-0012-1	177 g in 1 BOTTLE; Type 0: Not a Combination Product	02/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	02/17/2020	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 1/2026 Bayer HealthCare LLC.