TOLNAFTATE ANTIFUNGAL POWDER- tolnaftate powder
SCHOLL'S WELLNESS COMPANY LLC

Dr. Scholl's Tolnaftate Antifungal Powder

Active Ingredient

Tolnaftate 1%

Purpose

Antifungal

Use

- Clinically proven to prevent most athlete's foot (tinea pedis) 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.

Directios

- to prevent athlete's foot, wash and dry feet 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 change shoes and socks at least once daily

Other Information

Store between 20° to 25°C (68° to 77°F)

Inactive Ingredients

Chloroxylenol, Fragrance, Kaolin, Sodium Bicarbonate, Tricalcium Phosphate, Zea Mays (Corn) Starch, Zinc Oxide

Questions ?

1-866-360-3226

ANTIFUNGAL POWDER

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24-HOUR

ODOR PROTECTION

✓ Clinically proven to prevent most athlete's foot

✓ Helps stop recurrence with daily use

NET WT 4 OZ (113 g)

Drug Facts

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Dist. by: Scholl's Wellness Co., LLC
Parsippany, NJ 07054. Made in the USA
with globally sourced materials. C1523

LABSCH001024B-0

TOLNAFTATE ANTIFUNGAL POWDER

tolnaftate powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73469-0619
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Tolnaftate (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	Tolnaftate	1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q)	
KAOLIN (UNII: 24H4NWX5CO)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TRICALCIUM PHOSPHATE (UNII: K4C08XP666)	
STARCH, CORN (UNII: O8232NY3SJ)	
ZINC OXIDE (UNII: SOI2LOH54Z)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73469-0619-4	113 g in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	11/26/2025	

Labeler - SCHOLL'S WELLNESS COMPANY LLC (117174744)

Revised: 12/2025 SCHOLL'S WELLNESS COMPANY LLC