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

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### 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

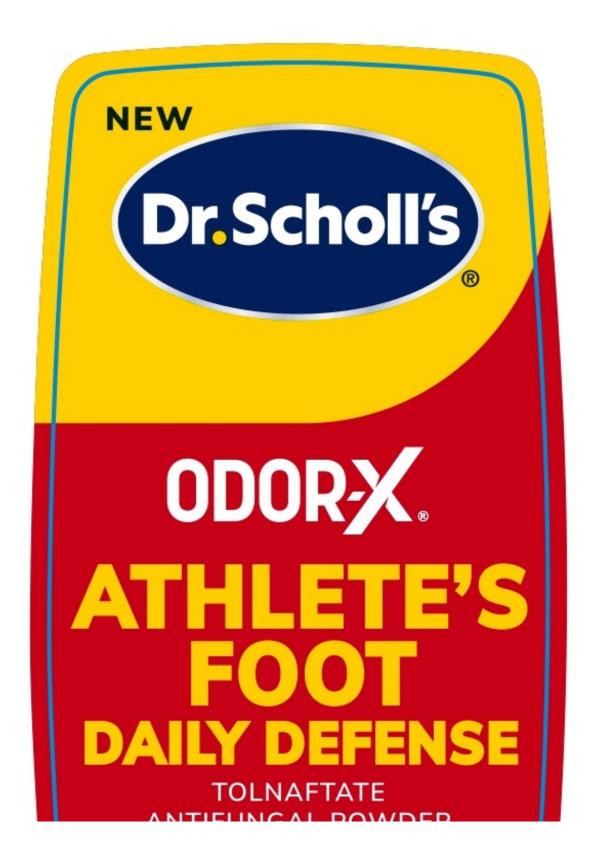
Store between 20° to 25°C (68° to 77°F)

### **Inactive Ingredients**

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

#### **Questions?**

1-866-360-3226



ANTIFUNGAL PUWDER

# + 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	<b>Basis of Strength</b>	Strength
П			

Tolnaftate (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV) Tolnaftate 1 g in 100 g

### **Inactive Ingredients**

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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**

#	tem 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

	<b>9</b>			
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