

**TINACTIN- tolnaftate cream**  
**Bayer HealthCare LLC.**

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**Tinactin ®**

***Drug Facts***

**Active ingredient**

Tolnaftate 1%

**Purpose**

Antifungal

**Uses**

- proven clinically effective in the treatment of most athlete's foot (tinea pedis) and ringworm (tinea corporis)
- helps prevent most athlete's foot with daily use
- for effective relief of itching, burning and cracking

**Warnings**

**For external use only**

**Do not use** on children under 2 years of age except under the advice and supervision of a doctor.

**When using this product** avoid contact with the eyes

**Stop use and ask a doctor if**

- irritation occurs
- there is no improvement within 4 weeks

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

**Directions**

- wash affected area and dry thoroughly
- apply a thin layer over affected area twice daily (morning and night)
- supervise children in the use of this product
- for athlete's foot: pay special attention to spaces between the toes; wear well-fitting, ventilated shoes and change shoes and socks at least once daily
- use daily for 4 weeks; if condition persists longer, ask a doctor
- to prevent athlete's foot, apply once or twice daily (morning and/or night)
- this product is not effective on the scalp or nails

**Other information**

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

**Inactive ingredients**

ceteth-20, cetostearyl alcohol, chlorocresol, mineral oil, propylene glycol, purified water, sodium phosphate monobasic, white petrolatum

**Questions?**

1-866-360-3266

Distributed by

Bayer HealthCare LLC, Whippany, NJ, USA, 07981

**PRINCIPAL DISPLAY PANEL - 15 g Tube Carton**



**TOUCH ACTIN'**

**Tinactin** ®

tolnaftate **ANTIFUNGAL**  
**CURES AND PREVENTS**  
**MOST ATHLETE'S FOOT**

**Relieves:**

- **itching**
- **burning**
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**CREAM**

# NET WT 15G (1/2 OZ)

## TINACTIN

tolnaftate cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11523-1190
<b>Route of Administration</b>	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
CETETH-20 (UNII: I835H2IHHX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CHLOROCRESOL (UNII: 36W5307109)	
MINERAL OIL (UNII: T5L8T28FGP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
PETROLATUM (UNII: 4T6H12BN9U)	

### Product Characteristics

<b>Color</b>	white (White to Off-white)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-1190-1	1 in 1 CARTON	12/12/2002	
1		15 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:11523-1190-2	1 in 1 CARTON	12/12/2002	
2		30 g in 1 TUBE; Type 0: Not a Combination Product		

### Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M005	09/23/1993	

**Labeler** - Bayer HealthCare LLC. (112117283)

Revised: 12/2025

Bayer HealthCare LLC.