

**CLOTRIMAZOLE- clotrimazole solution**  
**Sun Pharmaceutical Industries, Inc.**

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**Clotrimazole Topical Solution USP,1%**

***Drug Facts***

**Active ingredient**

Clotrimazole USP, 1%

**Purpose**

Antifungal

**Uses**

- cures most athlete's foot (tinea pedis), jock itch (tinea cruris), and ringworm (tinea corporis)
- effectively relieves
- itching
- cracking
- burning
- discomfort

which can accompany these conditions

**Warnings**

**For external use only**

**Ask a doctor before use**

- on children under 2 years of age

**When using this product**

- avoid contact with eyes

**Stop use and ask a doctor if**

- irritation occurs
- there is no improvement within 4 weeks (for athlete's foot and ringworm) or 2 weeks (for jock itch)

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

**Directions**

- This product is not effective on the scalp or nails For best results, follow directions and continue treatment for length of time indicated. For athlete's foot and ringworm:

use daily for 4 weeks. For jock itch: use daily for 2 weeks.

- clean the affected area and dry thoroughly
- apply a thin layer of the product over affected area twice daily (morning and night) or as directed by a doctor
- 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; change shoes and socks at least once daily

### Other information

- **Store at 20° to 25°C (68° to 77°F)**[see USP Controlled Room Temperature].

### Inactive ingredient

polyethylene glycol 400

### Questions?

Call **1-866-923-4914**

Distributed by:

**Taro Pharmaceuticals U.S.A., Inc.**

Hawthorne, NY 10532

### Clotrimazole Topical Solution USP 1%

Clotrimazole Topical Solution USP 1%

**NDC 51672-2158-1**

**Compare to the active ingredient in Lotrimin® AF\***

**Cures Most Athlete's Foot**

**Athlete's Foot Solution**

**Clotrimazole Topical Solution USP, 1%**

**Antifungal**

- Relieves Itching, Burning,
- Greaseless, Nonstaining

**10 mL (1/3 fl oz)**

\*All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Lotrimin® AF.

Distributed by:  
Taro Pharmaceuticals U.S.A., Inc.  
Hawthorne, NY 10532

Made in Canada  
TARO is a registered trademark of Taro Pharmaceuticals U.S.A., Inc.

51672-21581 1

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**Drug Facts**

Active ingredient	Purpose
Clotrimazole USP, 1%	Antifungal

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**Drug Facts (continued)**

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**Inactive ingredients**

polyethylene glycol 400

**Questions?**  
Call 1-866-923-4914

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**Clotrimazole Topical Solution 1%**

Compare to the  
active ingredient  
in Lotrimin<sup>®</sup> AF\*

*Cures Most Athlete's Foot*

# Athlete's Foot Solution

**Clotrimazole Topical Solution USP, 1%**

**Antifungal**

- *Relieves Itching & Burning*
- *Greaseless,  
Nonstaining*



**10 mL**  
**(1/3 fl oz)**

Cures Most  
Athlete's Foot

Antifungal

# Athlete's Foot Solution

- Relieves Itching & Burning
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Clotrimazole Topical Solution USP, 1%

## Drug Facts

### Active ingredient

Clotrimazole USP, 1%.....Antifungal

### Purpose

Antifungal

### Uses

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**Inactive ingredient** polyethylene glycol 400

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Hawthorne, NY 10532

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Taro Pharmaceuticals U.S.A., Inc.



TARO Pharmaceuticals USA, Inc.  
Made in Canada.  
PK-1627-4 0416-4

**TARO**

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## CLOTRIMAZOLE

clotrimazole solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:51672-2037
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CLOTRIMAZOLE</b> (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)	CLOTRIMAZOLE	1 g in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51672-2037-1	1 in 1 CARTON	05/01/1996	
1		10 mL in 1 BOTTLE; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	05/01/1996	

## CLOTRIMAZOLE

clotrimazole solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:51672-2158
<b>Route of Administration</b>	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CLOTRIMAZOLE (UNII: G07GZ97H65) (CLOTRIMAZOLE - UNII:G07GZ97H65)	CLOTRIMAZOLE	1 g in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51672-2158-1	1 in 1 CARTON	09/23/2024	
1		10 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	09/23/2024	

**Labeler** - Sun Pharmaceutical Industries, Inc. (146974886)

### Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(51672-2037, 51672-2158)

Revised: 7/2025

Sun Pharmaceutical Industries, Inc.