

**DRYELLO TOLNAFTATE 1% ANTIFUNGAL- tolnaftate stick**  
**ProPedix, Inc.**

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**Dryello Tolnaftate 1% Antifungal Improved (v1b) - 0.53 oz Stick - 2pk**

**Active Ingredient**

Tolnaftate 1%

**Purpose**

Antifungal

**Uses**

- Cures most Athlete's Foot (tinea pedis)
- Relieves itching, burning, cracking, and scaling
- Prevents recurrence of most Athlete's foot with daily use

**Warnings**

Do not use on children under 2 years of age unless directed by a doctor.

**For external use only.** Avoid contact with eyes.

**When using this product**

- If irritation occurs, discontinue use and consult a doctor

**Keep out of reach of children**

If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

- Wash feet and dry thoroughly.
- Apply a thin layer of the product to the affected area twice daily (morning and night).
- Supervise children in the use of this product.
- Pay special attention to spaces between toes; wear well-fitting, ventilated shoes, and change shoes and socks at least once daily.

**Other**

Store between 68° - 86°F (20° - 30°C)

**Inactive ingredients**

Aluminum Hydroxide, Arrowroot Powder, Behenyl Behenate, C12-15 Alkyl Benzoate, Cyclopentasiloxane, Magnesium Aluminum Silicate, Magnesium Citrate, Magnesium Hydroxide, Magnesium Sulfate, Orange Peel Oil, Stearyl Alcohol, Tocopheryl Acetate.

## Questions?

1-800-452-7362 dryello.com

## Principal Display Panel

For Athlete's Foot

Treats + Prevents

New Silky Smooth Application

Dryello™

TOLNAFTATE 1% ANTIFUNGAL

Clinically Proven

Dermatologist Developed

Two Glide-On Sticks

Two 15g (0.53oz) NET WT Sticks



**DRYELLO TOLNAFTATE 1% ANTIFUNGAL**

tolnaftate stick

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:85565-005
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
TOLNAFTATE (UNII: 06KB629TKV) (TOLNAFTATE - UNII:06KB629TKV)	TOLNAFTATE	1 g in 100 g

### Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: VR1WPI7EW8)	
C12-15 ALKYL BENZOATE (UNII: A9EJ3J61HQ)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
MARANTA ARUNDINACEA ROOT (UNII: FVN346W31A)	
MAGNESIUM HYDROXIDE (UNII: NBZ3QY004S)	
MAGNESIUM CITRATE (UNII: RHO26O1T9V)	
BEHENYL BEHENATE (UNII: K8NU647RJ0)	
ORANGE OIL (UNII: AKN3KSD11B)	
CYCLOPENTASILOXANE (UNII: 0THT5PCI0R)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
MAGNESIUM SULFATE (UNII: DE08037SAB)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85565-005-01	2 in 1 BOX	04/03/2026	
1		15 g in 1 APPLICATOR; 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	04/03/2026	

**Labeler** - ProPedix, Inc. (119483645)

### Establishment

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
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Revised: 4/2026

ProPedix, Inc.