

**DEFENSE ANTIFUNGAL MEDICATED BAR- tolnaftate soap**  
**Defense Soap LLC**

-----  
**Defense Antifungal Medicated Bar**

**Active ingredient**

Tolnaftate 1%

**Purpose**

Antifungal

**Uses**

- Proven clinically effective in the treatment of most athlete's foot (tinea pedis), ringworm (tinea corporis) and jock itch (tinea cruris).
- For effective relief of itching, burning and cracking skin.

**Warnings**

For external use only

**When using this product**

When using this product avoid contact with the eye

**Stop use and ask a doctor if**

***Stop use and ask a doctor if*** irritation occurs or if there is no improvement within 2 weeks for jock itch or within 4 weeks for athlete's foot and ringworm.

**Do not use**

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

**Keep out of reach of children.**

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

**Directions**

- Wash affected area well and let product sit 1-2 minutes before rinsing.
- Make sure to dry affected area thoroughly.
- Supervise children in the use of this product.
- For Athlete's foot: pay special attention to the spaces between the toes; wear well-fitting, ventilated shoes and change shoes and socks at least once daily.

- Use 1-2 times daily. Use daily for 4 weeks; if condition persists longer, ask a doctor.
- This product is not effective on the scalp or nails.

## Inactive ingredients

Sodium Palmate, Sodium Palm Kernelate, Water, Glycerin, Sodium Gluconate, Eucalyptus Globulus Leaf Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Palm Acid, Sodium Chloride, Palm Kernel Acid, Charcoal Powder

## Principal Display Panel

Tolnaftate 1%

Antifungal

Medicated Bar Soap

Cures and Prevents Most Fungal Skin Infections, Including Athlete's Foot, Ringworm and Jock Itch.



DEFENSE SOAP LLC, 3390 LIBERTY ATHENS AVE VERMILLION, OH 44089 866-544-1689

**DEFENSE ANTIFUNGAL MEDICATED BAR**

tolnaftate soap

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70856-3581
<b>Route of Administration</b>	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
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
WATER (UNII: 059QF0KO0R)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	
SODIUM PALM KERNELATE (UNII: 6H91L1NXTW)	
GLYCERIN (UNII: PDC6A3C0OX)	
MELALEUCA ALTERNIFOLIA LEAF (UNII: G43C57162K)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ACTIVATED CHARCOAL (UNII: 2P3VWUJ3H10)	
SODIUM PALMATE (UNII: S0A6004K3Z)	
SODIUM COCOATE (UNII: R1TQH25F4I)	

### Product Characteristics

<b>Color</b>	black	<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:70856-3581-3	119 g in 1 CARTON; Type 0: Not a Combination Product	02/01/2019	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	02/01/2019	

