AZOLEN- miconazole nitrate tincture Stratus Pharmaceuticals, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

AZOLEN™ TINCTURE

Drug Facts

Active Ingredient

Miconazole Nitrate 2%

Purpose

Topical Antifungal

Uses

- Cures most athlete's foot (tinea pedis) and ringworm (tinea corporis)
- For effective relief of itchy, scaly skin between the toes.

WARNINGS

- For external use only.
- Do not use on children under 2 years of age except under supervision of doctor.
- Avoid contact with eyes.

If irritation occurs or if there is no improvement within four weeks, discontinue use and consult a doctor.

Keep out of the reach of children. If swallowed, get medical help or contact poison control center immediately.

Directions

- Clean and dry the affected area.
- Cover the affected area with a thin layer twice a day (a.m. and p.m.) on skin, under nails and surrounding cuticle areas.
- Supervise children in the use of this product.
- For athlete's foot and ringworm use daily for four (4) weeks.
- This product is not effective on scalp or nails.

Other Information

• Store at 25°C (77°F); excursions permitted to 15°C - 30°C (59° - 86°F). [See USP Controlled Room Temperature]. Protect from freezing. If freezing occurs, warm to room temperature.

Inactive Ingredients

Benzyl Alcohol, Glacial Acetic Acid, Isopropyl Alcohol, Laureth-4, Purified Water and Sodium

Hydroxide Solution.

For more information, see enclosed package insert.

Questions?

1-800-442-7882

Distributed by: STRATUS PHARMACEUTICALS INC 12379 SW 130th Street Miami, Florida 33186

PRINCIPAL DISPLAY PANEL - 29.57 mL Bottle Carton

NDC 58980-818-10

 $\begin{array}{c} \textbf{AZOLEN}^{\text{TM}} \\ \textbf{TINCTURE} \end{array}$

(MICONAZOLE NITRATE USP, 2%)

TOPICAL ANTIFUNGAL

Distributed by: **STRATUS**

NET VOL.: 1.0 FL OZ (29.57 mL)



NET VOL : 1.0 FL OZ (29.57 mL)

(MICONAZOLE NITRATE USP, 2%)

TINCTURE



NDC 28980-818-10

NDC 58980-818-10



ZOLEN

TINCTURE

(MICONAZOLE NITRATE USP, 2%)

TOPICAL ANTIFUNGAL

Distributed by:



NET VOL.: 1.0 FL OZ (29.57 mL)

Drug Facts
Active Ingredient
Miconazole Nitrate 2%
Purpose
Topical Antifungal

Uses

 Cures most athlete's foot (tinea pedis) and ringworm (tinea corporis)
 For effective relief of itchy, scaly skin between the toes.

WARNINGS ● For external use only. ● Do not use on children under 2 years of age except under supervision of doctor. ● Avoid contact with eyes.

If irritation occurs or if there is no improvement within four weeks, discontinue use and consult a doctor.

Keep out of the reach of children. If swallowed, get medical help or contact poison control center immediately.

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NDC 58980-818-10



TINCTURE

(MICONAZOLE NITRATE USP, 2%)

TOPICAL ANTIFUNGAL

Distributed by:



NET VOL.: 1.0 FL OZ (29.57 mL)

Drug Facts (continued)

Directions

· Clean and dry the affected area. . Cover the affected area with a thin layer twice a day (a.m. and p.m.) on skin, under nails and surrounding cuticle areas. . Supervise children in the use of this product. . For athlete's foot and ringworm use daily for four (4) weeks. • This product is not effective on scalp or nails.

Other Information . Store at 20°C to 25°C (68°F to 77°F), excursions permitted between 15°C and 30°C (between 59°F and 86°F). [See USP Controlled Room Temperature]. Protect from freezing. If freezing occurs, warm to room temperature.

Inactive Ingredients: Benzyl Alcohol, Glacial Acetic Acid, Isopropyl Alcohol, Laureth-4, Purified Water and Sodium Hydroxide Solution.

For more information, see enclosed package insert.

Questions?: 1-800-442-7882

VC-AT1-2012-340









PLEASE LEAVE UNVARNISHED. AREA FOR EXP DATE & LOT



AZOLEN

miconazole nitrate tincture

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Product TypeHUMAN OTC DRUG LABELItem Code (Source)NDC:58980-818

Route of Administration TOPICAL DEA Schedule

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
Miconazole Nitrate (Miconazole)

Miconazole Nitrate
20 mg in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
Water				
Benzyl Alcohol				
Acetic Acid				
Isopropyl Alcohol				
Laureth-4				
Sodium Hydroxide				

]	Packaging					
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:58980-818-10	1 in 1 BOX				
1	L	29.57 mL in 1 BOTTLE, WITH APPLICATOR				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH NOT FINAL	part333C	12/0 1/20 11		

Labeler - Stratus Pharmaceuticals, Inc. (789001641)

Establishment					
Name	Address	ID/FEI	Business Operations		
Sonar Products Inc		104283945	MANUFACTURE(58980-818)		

Revised: 2/2013 Stratus Pharmaceuticals, Inc.