

**TRIPLE ANTIFUNGAL- miconazole nitrate ointment**  
**Advantice Health, LLC**

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**Triple Paste Antifungal Ointment**

***Drug Facts***

**Active ingredient**

Miconazole nitrate 2%

**Purpose**

Antifungal

**Uses**

- For the treatment of superficial skin infections caused by yeast (*Candida albicans*)
- Relieves discomfort, irritation, redness and chafing associated with jock itch
- Cures most jock itch (tinea cruris)

**Warnings**

**For external use only**

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

**Stop use and ask a doctor if**

- irritation occurs
- there is no improvement within 2 weeks

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

**Keep out of reach of children.**

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

**Directions**

- Clean affected area and dry thoroughly
- Apply a thin layer of product over affected area twice daily (morning and night) or as directed by a doctor
- Supervise children in the use of this product
- Use daily for 2 weeks; If condition persists longer, consult a doctor
- This product is not effective on the scalp or nails

**Other information**

Store at room temperature 15°- 30°C (59°- 86°F)

## **Inactive ingredients**

White petrolatum, zinc oxide, corn starch, anhydrous lanolin, stearyl alcohol, beeswax, cholesterol.

## **Questions?**

1-800-345-0032

Distributed by Advantice Health, LLC  
Bridgewater, NJ 08807

## **PRINCIPAL DISPLAY PANEL - 56.7 g Tube Carton**

Triple Paste AF <sup>®</sup>

2% MICONAZOLE NITRATE ANTIFUNGAL OINTMENT

- Relieves burning, redness, irritation and discomfort
- Unique, soothing, water-resistant formula
- Lubricates skin to heal and prevent chafing

Help is on the way...

Net Wt. 2 oz (56.7 g)



## TRIPLE ANTIFUNGAL

miconazole nitrate ointment

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:16864-040
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII: 7NNO0D7S5M)	MICONAZOLE NITRATE	2 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>WHITE PETROLATUM</b> (UNII: B6E5W8RQJ4)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>LANOLIN</b> (UNII: 7EV65EAW6H)	
<b>STEARYL ALCOHOL</b> (UNII: 2KR89I4H1Y)	
<b>YELLOW WAX</b> (UNII: 2ZA36H0S2V)	
<b>LEVOMENOL</b> (UNII: 24WE03BX2T)	
<b>CHOLESTEROL</b> (UNII: 97C5T2UQ7J)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>OAT</b> (UNII: Z6J799EAJK)	

## Product Characteristics

<b>Color</b>	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:16864-040-02	1 in 1 CARTON	09/01/2021	
1		56.7 g in 1 TUBE; 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	09/01/2021	

**Labeler** - Advantice Health, LLC (192527062)

Revised: 12/2024

Advantice Health, LLC