

# **MICONAZOLE NITRATE- miconazole nitrate cream**

## **A-S Medication Solutions**

-----

### **Miconazole Nitrate**

#### ***Drug Facts***

#### **Active ingredient**

Miconazole nitrate USP 2% (100 mg in each applicatorful)

#### **Purpose**

Vaginal antifungal

#### **Uses**

- treats vaginal yeast infections
- relieves external itching and irritation due to a vaginal yeast infection

#### **Warnings**

##### **For vaginal use only**

**Do not use if you have never had a vaginal yeast infection diagnosed by a doctor**

##### **Ask a doctor before use if you have**

- **vaginal itching and discomfort for the first time**
- **lower abdominal, back or shoulder pain, fever, chills, nausea, vomiting, or foul-smelling vaginal discharge. You may have a more serious condition.**
- vaginal yeast infections often (such as once a month or 3 in 6 months). You could be pregnant or have a serious underlying medical cause for your symptoms, including diabetes or a weakened immune system.
- been exposed to the human immunodeficiency virus (HIV) that causes AIDS

**Ask a doctor or pharmacist before use if you are** taking the prescription blood thinning medicine warfarin, because bleeding or bruising may occur

##### **When using this product**

- do not use tampons, douches, spermicides, or other vaginal products. Condoms and diaphragms may be damaged and fail to prevent pregnancy or sexually transmitted diseases (STDs).
- do not have vaginal intercourse
- mild increase in vaginal burning, itching or irritation may occur
- if you do not get complete relief ask a doctor before using another product

**Stop use and ask a doctor if**

- **symptoms do not get better in 3 days**
- **symptoms last more than 7 days**
- **you get a rash or hives, abdominal pain, fever, chills, nausea, vomiting, or foul-smelling vaginal discharge.**

**If pregnant or breast-feeding,** ask a health professional before use.

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

### **Directions**

- before using this product read the enclosed consumer information leaflet for complete directions and information
- adults and children 12 years of age and over:
  - **applicator:** insert 1 applicatorful into the vagina at bedtime for 7 nights in a row. Wash applicator after use.
  - use the same tube of cream if you have itching and irritation on the skin outside the vagina. Squeeze a small amount of cream onto your fingertip. Apply to itchy, irritated skin outside the vagina (vulva). Use 2 times daily for up to 7 days as needed.
- **children under 12 years of age: ask a doctor**

### **Other information**

- do not purchase if carton is opened
- do not use if seal over tube opening has been punctured or is not visible
- store at 20° to 25°C (68° to 77°F)
- see carton or tube crimp for lot number and expiration date

### **Inactive ingredients**

benzoic acid (0.20%) as a preservative, butylated hydroxytoluene, mineral oil, oleoyl polyoxyglyceride, PEG-6-32 stearate/glycol stearate, and purified water.

### **Questions?**

call **1-866-923-4914**

Distributed by:

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

Hawthorne, NY 10532

### **HOW SUPPLIED**

Product: 50090-4356

NDC: 50090-4356-0 45 g in a TUBE, WITH APPLICATOR / 1 in a CARTON

# Miconazole Nitrate



## MICONAZOLE NITRATE

miconazole nitrate cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50090-4356(NDC:51672-2035)
<b>Route of Administration</b>	VAGINAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>MICONAZOLE NITRATE</b> (UNII: VV4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	20 mg in 1 g

### Inactive Ingredients

Ingredient Name	Strength
<b>Butylated Hydroxytoluene</b> (UNII: 1P9D0Z171K)	
<b>Mineral Oil</b> (UNII: T5L8T28FGP)	
<b>Water</b> (UNII: 059QF0KO0R)	
<b>Benzoic Acid</b> (UNII: 8SKN0B0MIM)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-4356-0	1 in 1 CARTON	06/14/2019	
1		45 g in 1 TUBE, WTH APPLICATOR; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA074444	01/13/1997	

**Labeler** - A-S Medication Solutions (830016429)

## Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-4356)

Revised: 12/2023

A-S Medication Solutions