

MICONAZOLE 7- miconazole nitrate cream
NuCare Pharmaceuticals, Inc.

Miconazole Nitrate Vaginal Cream, USP

Drug Facts

Active ingredient

Miconazole nitrate, USP 2% (100 mg in each applicator)

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 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

- to open tube use cap to puncture seal
- do not use if seal over tube opening has been punctured or is not visible
- do not purchase if carton is open
- store at 20°-25°C (68°-77°F) [see USP Controlled Room Temperature] excursions permitted to 15°-30°C (59°-86°F).

Inactive ingredients

benzoic acid, butylated hydroxyanisole, mineral oil, oleoyl polyoxyglycerides, pegoxol 7 stearate, purified water

Questions?

1-800-432-8534 between 9 am and 4 pm EST, Monday-Friday.

Principal display panel

NuCare Pharmaceuticals, Inc.

NDC: 68071-1703-7
Miconazole 7 2%
45g Vaginal Cr.
 Miconazole Nitrate, USP 2%
 (100mg in each applicator)
 See manufacturer's label
 for full list of ingredients.

Product #: R0299045

Warnings:
 WARNING: KEEP OUT OF REACH OF CHILDREN
 STORE AT CONTROLLED TEMPERATURE 59-86°F.

Lot and Expiration Information:
 Miconazole 7 2%
 Lot: 000000 NDC: 68071-1703-07
 MFR NDC: 0472-0730-41 Exp.: 00-00
 Serial# 00000000002

GTIN and Serial Information:
 GTIN 00368071170375
 Serial# 000000000002
 Exp. Date 00-00
 LOT#: 000000

Manufacturer and Distribution:
 Distributed by: Actavis Pharma, Inc. Parsippany, NJ 07054
 Packaged By: NuCare Pharmaceuticals, Inc. Orange, CA 92867

Barcode and Patient Instructions:
 6807117030745000000000000
 Use only as directed by your physician.

MICONAZOLE 7			
miconazole nitrate cream			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-1703(NDC:0472-0730)
Route of Administration	VAGINAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
MICONAZOLE NITRATE (UNII: VW4H1CYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)		MICONAZOLE NITRATE	20 mg in 1 g
Inactive Ingredients			
Ingredient Name			Strength
BENZOIC ACID (UNII: 8SKN0B0MIM)			
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)			
MINERAL OIL (UNII: T5L8T28FGP)			
APRICOT KERNEL OIL PEG-6 ESTERS (UNII: DRG3KJZ1TJ)			
PEGOXOL 7 STEARATE (UNII: 3EW5AXE5X5)			
WATER (UNII: 059QF0KO0R)			

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-1703-7	45 g in 1 BOX; Type 0: Not a Combination Product	03/04/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA074164	04/01/1997	

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-1703)

Revised: 1/2022

NuCare Pharmaceuticals, Inc.