

MICONAZOLE 3 COMBINATION PACK- miconazole nitrate
Major Pharmaceuticals

Miconazole 3 Combination Pack

Drug Facts

Active ingredients

Miconazole nitrate, USP 200 mg (in each suppository)
Miconazole nitrate, USP 2% (external cream)

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 and 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.

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 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:**
 - **suppositories:** insert 1 suppository into the vagina at bedtime for 3 nights in a row. Throw applicator away after use.
 - **external cream:** Use cap to puncture seal. Squeeze a small amount of cream onto your fingertip. Apply the cream onto the itchy, irritated skin outside the vagina. Use 2 times daily for up to 7 days, as needed.
- **children under 12 years of age: ask a doctor**

Other information

- do not use if suppository wrapper is missing or damaged (each suppository is individually wrapped)
- 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

- suppository: hydrogenated vegetable oil
- external cream: benzoic acid, butylated hydroxyanisole, mineral oil, oleoyl polyoxyglycerides, pegoxol 7 stearate, purified water

Questions?

1-888-838-2872 between 9 am and 5 pm ET, Monday-Friday.

Principal Display Panel

MAJOR[®]

NDC 0904-5415-01

MICONAZOLE 3

MICONAZOLE NITRATE VAGINAL SUPPOSITORIES, USP (200 MG)

& MICONAZOLE NITRATE VAGINAL CREAM, USP 2% VAGINAL ANTIFUNGAL

COMBINATION PACK

CURES MOST VAGINAL YEAST INFECTIONS

AND RELIEVES ASSOCIATED EXTERNAL ITCHING & IRRITATION

COMPARE TO the active ingredient in MONISTAT[®] 3*

3 MICONAZOLE NITRATE

VAGINAL SUPPOSITORIES,

USP 200 mg PLUS 1 TUBE OF

MICONAZOLE NITRATE VAGINAL
CREAM, USP 2%

NET WT. 0.32 OZ (9g) TUBE

MAJOR®

MICONAZOLE 3

MICONAZOLE NITRATE VAGINAL SUPPOSITORIES, USP (200 mg) & MICONAZOLE NITRATE VAGINAL CREAM, USP 2% VAGINAL ANTIFUNGAL

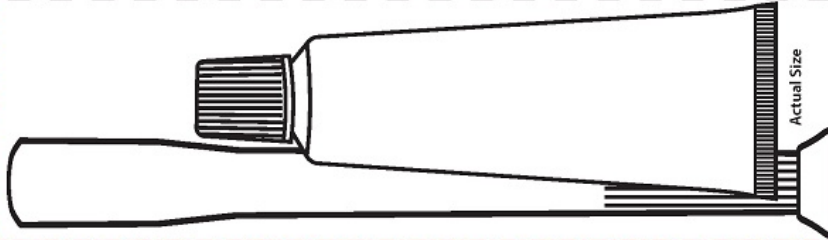
COMBINATION PACK

3 DAY TREATMENT



3 Vaginal Suppositories with Disposable Applicators

PLUS EXTERNAL CREAM FOR ITCH RELIEF



MAJOR®

NDC 0904-5415-01

MICONAZOLE 3

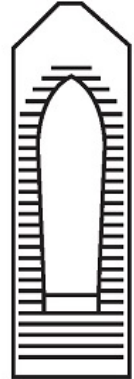
MICONAZOLE NITRATE VAGINAL SUPPOSITORIES, USP (200 mg) & MICONAZOLE NITRATE VAGINAL CREAM, USP 2% VAGINAL ANTIFUNGAL

COMBINATION PACK

CURES MOST VAGINAL YEAST INFECTIONS AND RELIEVES ASSOCIATED EXTERNAL ITCHING & IRRITATION

3 MICONAZOLE NITRATE VAGINAL SUPPOSITORIES, USP 200 mg PLUS 1 TUBE OF MICONAZOLE NITRATE VAGINAL CREAM, USP 2% NET WT. 0.32 OZ (9g) TUBE

3 Suppositories



COMPARE TO the active ingredient in MONISTAT® 3*

*This product is not affiliated with, manufactured by, or produced by the makers or owners of Monistat®.

73100618 R0219
GW710208

Made in USA

Distributed by: MAJOR® PHARMACEUTICALS

17177 N Laurel Park Dr., Suite 233

Livonia, MI 48152 USA

Rev. 02/19 Re-Order No. 700600 M-46



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Suppositories: Insert 1 suppository into the vagina at bedtime for 3 nights in a row. Throw applicator away after use.

Adults and children 12 years of age and over:

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

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
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Stop use and ask a doctor if:

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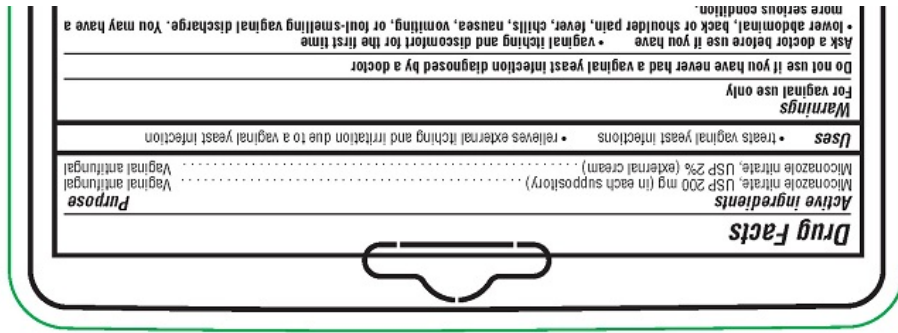
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MICONAZOLE 3 COMBINATION PACK

miconazole nitrate kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0904-5415
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-5415-01	1 in 1 PACKAGE, COMBINATION; Type 0: Not a Combination Product	12/02/2002	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BLISTER PACK	3 in 0
Part 2	1 TUBE	9 g

Part 1 of 2

MICONAZOLE 3

miconazole nitrate suppository

Product Information

Route of Administration	VAGINAL
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Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MICONAZOLE NITRATE (UNII: VW4HICYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	200 mg

Inactive Ingredients

Ingredient Name	Strength
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HYDROGENATED PALM KERNEL OIL (UNII: FM8D1RE2VP)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		3 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA074926	12/02/2002	

Part 2 of 2

MICONAZOLE 3

miconazole nitrate cream

Product Information

Route of Administration VAGINAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MICONAZOLE NITRATE (UNII: VW4HICYW1K) (MICONAZOLE - UNII:7NNO0D7S5M)	MICONAZOLE NITRATE	10 mg in 0.5 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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		9 g in 1 TUBE; Type 0: Not a Combination Product		

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA074926	12/02/2002	

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ANDA	ANDA074926	12/02/2002	

Labeler - Major Pharmaceuticals (191427277)

Revised: 11/2019

Major Pharmaceuticals