

MICONAZOLE 3- miconazole nitrate cream
TARGET Corporation

Miconazole 3

Drug Facts

Active ingredient

Miconazole nitrate USP 4%

Purpose

Vaginal antifungal

Use

- treats vaginal yeast infections

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:
 - insert 1 applicatorful into the vagina at bedtime for 3 nights in a row.
Throw applicator away after use.
- children under 12 years of age: ask a doctor

Other information

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

Inactive ingredients

cetyl alcohol, isopropyl myristate, polysorbate 60, polysorbate 80, potassium hydroxide, propylene glycol, purified water, sorbitan monostearate and stearyl alcohol.

Questions?

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

Dist. by Target Corp., Mpls., MN 55403

PRINCIPAL DISPLAY PANEL - 25 g Tube Carton

NDC 11673-391-06

Compare to the active ingredient in Monistat[®] 3*

up&up

3-day treatment

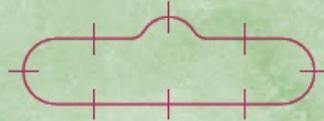
miconazole 3

miconazole nitrate vaginal cream USP (4%)

vaginal antifungal

cures most vaginal yeast infections

ONE 0.9 OZ (25 g) TUBE OF 3 DAY VAGINAL CREAM AND THREE DISPOSABLE APPLICATORS



NDC 11673-391-06

Compare to the active ingredient in Monistat® 3*

3-day treatment



miconazole 3

**miconazole nitrate vaginal cream USP (4%)
vaginal antifungal**

cures most vaginal yeast infections

ONE 0.9 OZ (25 g) TUBE OF 3 DAY VAGINAL CREAM AND THREE DISPOSABLE APPLICATORS

*All trademarks are property of their respective owners.
†This product is not affiliated with the makers/owners of Monistat® 3.

TAMPER EVIDENT:
DO NOT USE IF THE SEAL ON THE
TUBE IS PUNCTURED OR NOT VISIBLE

100% satisfaction guaranteed or your money back.

We welcome any questions you may have at
Target.com/comments or 1-800-910-6874.



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Other information

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- children under 12 years of age: ask a doctor.

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Questions? call 1-866-923-4914

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• been exposed to the human immunodeficiency virus (HIV) that causes AIDS

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- if you do not get complete relief ask a doctor before using another product

Drug Facts

Active ingredient
Micronazole nitrate USP 4%
Vaginal antifungal

Purpose
Vaginal antifungal

Use • treats vaginal yeast infections

Warnings

Do not use if you have never had a vaginal yeast infection diagnosed by a doctor.
For vaginal use only

Ask a doctor before use if you have

- lower abdominal, back or shoulder pain, fever, chills, nausea, vomiting or
- vaginal itching and discomfort for the first time

• lower abdominal vaginal discharge. You may have a more severe condition

Drug Facts (continued)

Stop use and ask a doctor if:

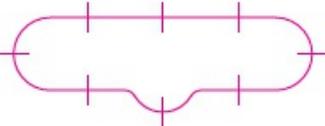
- symptoms do not get better in 3 days
- symptoms last more than 7 days
- you get a rash or hives, abdominal pain, fever, chills, nausea, vomiting or
- you get a foul-smelling vaginal discharge

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

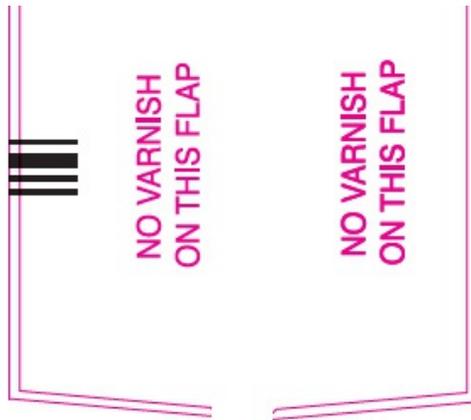
- Before using this product read the enclosed consumer information leaflet for complete directions and information
- adults and children 12 years of age and over.



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NO VARNISH/ NO AQ
NO COPY / NO COLOR
THIS FLAP FOR LOT #
AND EXP DATE PRINT

T128B
B81.3
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MICONAZOLE 3

miconazole nitrate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-391
Route of Administration	VAGINAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Miconazole Nitrate (UNII: VW4H1CYW1K) (Miconazole - UNII:7NNO0D7S5M)	Miconazole Nitrate	40 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
cetyl alcohol (UNII: 936JST6JCN)	
isopropyl myristate (UNII: 0RE8K4LNJS)	
polysorbate 60 (UNII: CAL22UVI4M)	
polysorbate 80 (UNII: 6OZP39ZG8H)	
potassium hydroxide (UNII: WZH3C48M4T)	
propylene glycol (UNII: 6DC9Q167V3)	
water (UNII: 059QF0K00R)	
sorbitan monostearate (UNII: NVZ4I0H58X)	
stearyl alcohol (UNII: 2KR89I4H1Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-391-06	1 in 1 CARTON	10/14/2020	
1		25 g in 1 TUBE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076773	03/02/2005	

Labeler - TARGET Corporation (006961700)

Revised: 10/2020

TARGET Corporation