MONISTAT 3 COMBINATION PACK- miconazole nitrate Insight Pharmaceuticals

MONISTAT®3 COMBINATION PACK

Miconazole Nitrate Vaginal Suppositories (200 mg) and Miconazole Nitrate Vaginal Cream (2%) Vaginal Antifungal

DRUG FACTS

Active ingredients	Purpose
Miconazole nitrate (200 mg in each suppository)	Vaginal antifungal
Miconazole nitrate 2% (external cream)	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:
- **suppositories:** insert 1 suppository into the vagina at bedtime for 3 nights in a row. Throw away applicator after use.
- **external cream:** 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 printed suppository blister is torn, open or incompletely sealed
- do not use if seal over tube opening has been punctured or embossed design \square is not visible
- do not purchase if carton is open
- store at 20°-25°C (68°-77°F)

Inactive ingredients

suppository: hydrogenated vegetable oil base

external cream: benzoic acid, cetyl alcohol, isopropyl myristate, polysorbate 60, potassium hydroxide, propylene glycol, purified water, stearyl alcohol

Questions? If you have any questions or comments, please call 1-877-666-4782 Visit our website @ www.monistat.com

PRINCIPAL DISPLAY PANEL - Kit Carton

3 DAY TREATMENT • SOFT INSERTS

MONISTAT[®] 3 Miconazole Nitrate Vaginal Suppositories (200 mg) and Miconazole Nitrate Cream (2%) (Miconazole Nitrate 200 mg per suppository plus 2% external vulvar cream) VAGINAL ANTIFUNGAL

CURES MOST VAGINAL YEAST INFECTIONS

and relieves associated external itching and irritation

CLINICALLY PROVEN MULTI-SYMPTOM RELIEF

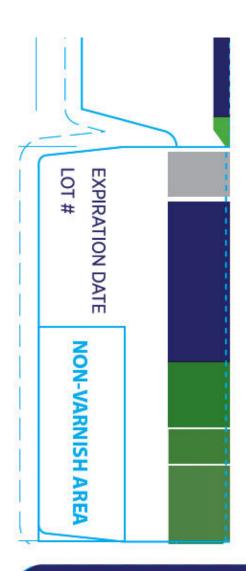
COMBINATION PACK

Net Wt. 3 Suppositories • 0.32 oz. (9g) tube



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DPEN HERE FUNGAL

MONISTAT³

Miconazole Nitrate Vaginal Suppositories (200 mg) and Miconazole Nitrate Cream (2%) VAGINAL ANTIFUNGAL

COMBINATION PACK

Drug Facts

Active ingredients

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 - (HIV) that causes AIDS

Drug Facts (continued)

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

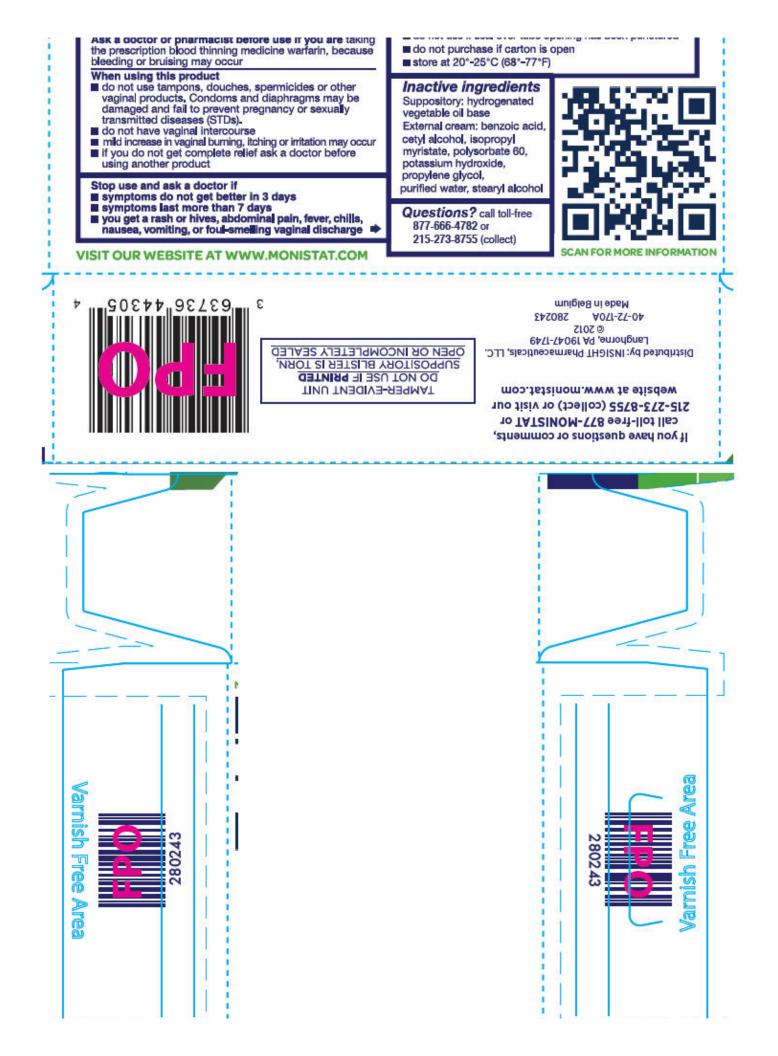
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Other information

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- do not use if seal over tube opening has been punctured







MONISTAT 3 CO miconazole nitrate kit	MBIN A	ATION PACK						
Product Information								
		DTC DRUG	Tearra	Code (Service)		NDC:63736-0	16	
Product Type	HUMAN C	JIC DRUG	Ite m	Code (Source)		NDC:03/30-0	10	
Packaging								
# Item Code	Ра	ckage Description		Marketing Start	Date	Marketing	End Date	
1 NDC:63736-016-03		KAGE, COMBINATION						
Quantity of Parts								
- 0	ckage Qua	antity		Total P	roduct Q	uantity		
Part 1 1 BLISTER PACK	0 0	5	3			. 0		
Part 2 1 TUBE			9 g					
Part 1 of 2								
MONISTAT 3	· · ·							
miconazole nitrate supp	ository							
Product Information								
Route of Administration		VAGINAL						
Active Ingredient/Ac	Active Ingredient/Active Moiety							
Ingredient Name				Basis o	of Strength	Strength		
Miconazole Nitrate (UNII: VW4H1CYW1K) (Miconazole - UNII:7NNO0D7S5M) Miconazole Nitrate			le Nitrate	200 mg				
Inactive Ingredients								
Ingredient Name Strength								
HYDROGENATED COCONUT OIL (UNII: JY810XM10M)								
HYDROGENATED PALM KERNEL OIL (UNII: FM8D1RE2VP)								
Packaging								
# Item Code	Package	Description	Ma	rketing Start Date		Marketing E	nd Date	
	0	•		5				

1	3 in 1 BLISTER	РАСК				
Marketing In	formation					
Marketing Catego	ry Applicati	on Number or Mon	ograph Citation	Marketing Start	t Date Marke	ting End Date
NDA	NDA020670			04/16/1996		
Part 2 of 2						
MONISTAT	3					
miconazole nitrate						
Product Inform	ation					
Route of Administ	ration	VAGINAL				
Active Ingredie	nt/Active Mo	ie ty				
	Ing	redient Name		Basis	of Strength	Strength
Miconazole Nitrate	(UNII: VW4H1CYV	W1K) (Miconazole - U	NII:7NNO0D7S5M)	Miconaz	ole Nitrate	10 mg in 0.5 g
The stars Terried						
Inactive Ingred	lenus	Ingredient Nar	na		St	rength
benzoic acid (UNII:	8 S K N 0 B 0 MIM)	Ingreutent Nai	IIC			Iengtii
cetyl alcohol (UNII:						
isopropyl myristate	(UNII: 0 RE8 K4LM	JS)				
polysorbate 60 (UN	II: CAL22UVI4M)					
propylene glycol (U		3)				
water (UNII: 059QF0						
water (UNII: 059QF0 stearyl alcohol (UN						
stearyl alcohol (UN						
stearyl alcohol (UN Packaging	II: 2KR8914H1Y)	Description	Marketing S	Start Date	Marketing	End Date
stearyl alcohol (UN	II: 2KR89I4H1Y) Package	Description	Marketing S	Start Date	Marketing	End Date
stearyl alcohol (UN Packaging # Item Code	II: 2KR8914H1Y)	Description	Marketing S	Start Date	Marketing	End Date
stearyl alcohol (UN Packaging # Item Code	II: 2KR89I4H1Y) Package	Description	Marketing S	Start Date	Marketing	End Date
stearyl alcohol (UN Packaging # Item Code 1	II: 2KR89I4H1Y) Package 9 g in 1 TUBE	Description	Marketing S	Start Date	Marketing	End Date
stearyl alcohol (UN Packaging # Item Code	E: 2KR89I4H1Y) Package 9 g in 1 TUBE formation	Description on Number or Mon		Start Date Marketing Start		End Date
stearyl alcohol (UN Packaging # Item Code 1 Marketing In	E: 2KR89I4H1Y) Package 9 g in 1 TUBE formation					
stearyl alcohol (UN Packaging # Item Code 1 Marketing In Marketing Catego	E: 2KR89I4H1Y) Package 9 g in 1 TUBE formation ry Applicati			Marketing Start		
stearyl alcohol (UN Packaging # Item Code 1 Marketing In Marketing Catego	E: 2KR89I4H1Y) Package 9 g in 1 TUBE formation ry Applicati			Marketing Start		

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA020670	04/16/1996	

Labeler - Insight Pharmaceuticals (176792315)

Establishment

Name	Address	ID/FEI	Business Operations
Jubilant		246762764	MANUFACTURE(63736-016)

Revised: 8/2012

Insight Pharmaceuticals