#### NYSTATIN- nystatin suspension Preferred Pharmaceuticals, Inc.

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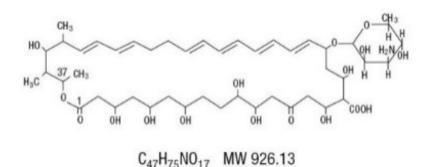
#### NYSTATIN ORAL SUSPENSION, USP

(100,000 units per mL)

**Rx only** 

## DESCRIPTION

Nystatin is an antimycotic polyene antibiotic obtained from *Streptomyces noursei*. Structural formula:



Nystatin Oral Suspension, for oral administration, contains 100,000 USP Nystatin Units per mL. Inactive ingredients: alcohol ( $\leq 1\%$  v/v), artificial wild cherry flavor, banana flavor, D&C yellow #10, FD&C red #40, glycerin, USP, magnesium aluminum silicate, methylparaben, NF, potassium phosphate dibasic, USP, propylene glycol, USP, propylparaben, NF, purified water, USP and sucrose 33.5%. May also contain citric acid, USP for pH adjustment.

# CLINICAL PHARMACOLOGY

### Pharmacokinetics

Gastrointestinal absorption of nystatin is insignificant. Most orally administered nystatin is passed unchanged in the stool. In patients with renal insufficiency receiving oral therapy with conventional dosage forms, significant plasma concentrations of nystatin may occasionally occur.

## Microbiology

Nystatin is both fungistatic and fungicidal *in vitro* against a wide variety of yeasts and yeast-like fungi. *Candida albicans* demonstrates no significant resistance to nystatin *in vitro* on repeated subculture in increasing levels of nystatin; other *Candida* species become quite resistant. Generally, resistance does not develop *in vivo*. Nystatin acts by binding to sterols in the cell membrane of susceptible *Candida* species with a resultant

change in membrane permeability allowing leakage of intracellular components. Nystatin exhibits no appreciable activity against bacteria, protozoa, or viruses.

### INDICATIONS AND USAGE

Nystatin Oral Suspension is indicated for the treatment of candidiasis in the oral cavity.

### CONTRAINDICATIONS

The preparation is contraindicated in patients with a history of hypersensitivity to any of its components.

### PRECAUTIONS

#### General

This medication is not to be used for the treatment of systemic mycoses. Discontinue treatment if sensitization or irritation is reported during use.

#### Carcinogenesis, Mutagenesis, Impairment of Fertility

No long-term animal studies have been performed to evaluate carcinogenic potential. There also have been no studies to determine mutagenicity or whether this medication affects fertility in males or females.

### Pregnancy

### **Teratogenic Effects Category C**

Animal reproduction studies have not been conducted with nystatin oral suspension. It is also not known whether nystatin oral suspension can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. Nystatin oral suspension should be given to a pregnant woman only if clearly needed.

#### **Nursing Mothers**

It is not known whether nystatin is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when nystatin is administered to a nursing woman.

### **Pediatric Use**

### See DOSAGE AND ADMINISTRATION.

### **ADVERSE REACTIONS**

Nystatin is well tolerated even with prolonged therapy. Oral irritation and sensitization have been reported. (See **PRECAUTIONS: General**).

*Gastrointestinal:* Diarrhea (including one case of bloody diarrhea), nausea, vomiting, gastrointestinal upset/disturbances.

*Dermatologic:* Rash, including urticaria has been reported rarely. Stevens-Johnson syndrome has been reported very rarely.

*Other:* Tachycardia, bronchospasm, facial swelling, and non-specific myalgia have also been rarely reported.

## OVERDOSAGE

Oral doses of nystatin in excess of five million units daily have caused nausea and gastrointestinal upset. There have been no reports of serious toxic effects of superinfections (see **CLINICAL PHARMACOLOGY, Pharmacokinetics**).

## DOSAGE AND ADMINISTRATION

INFANTS: 2 mL (200,000 units) four times daily (in infants and young children, use dropper to place one-half of dose in each side of mouth and avoid feeding for 5 to 10 minutes).

NOTE: Limited clinical studies in premature and low birth weight infants indicate that 1 mL four times daily is effective.

CHILDREN AND ADULTS: 4–6 mL (400,000 to 600,000 units) four times daily (one-half of dose in each side of mouth). The preparation should be retained in the mouth as long as possible before swallowing.

Continue treatment for at least 48 hours after perioral symptoms have disappeared and cultures demonstrate eradication of *Candida albicans*.

## HOW SUPPLIED

Nystatin Oral Suspension, USP, 100,000 USP Nystatin Units per mL, is available as a fruit flavored, light creamy yellow, ready-to-use suspension.

60 mL bottles was a calibrated dropper and

1 Pint (473 mL) bottles (68788-6792-6)

## Storage

Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Avoid freezing

## **Rx Only**

Product No.: 8537

Manufactured For:

Wockhardt USA, LLC

Parsippany, NJ 07054

Manufactured By:

Morton Grove Pharmaceuticals, Inc.

### Morton Grove, IL 60053

A50-8537-16

REV. 07-18

### Relabeled By: Preferred Pharmaceuticals Inc.

### **PRINCIPAL DISPLAY PANEL Bottle Carton**

#### MGP

NDC 68788-6792-6

#### NYSTATIN ORAL

SUSPENSION, USP

(100,000 units

per mL)

**Fruit Flavored** 

SHAKE WELL BEFORE USING

DO NOT USE IF INNER FOIL SEAL PRINTED "SEALED FOR YOUR PROTECTION" IS BROKEN OR MISSING

**Rx Only** 

NET: 1 Pint (473 mL)

### **Relabeled By: Preferred Pharmaceuticals Inc.**



Nystatin Label

## NYSTATIN

nystatin suspension

#### **Product Information**

Broduct Type							
Product Type		HUMAN PRESCRIPTION DRUG	ltem Co (Source		NDC:68 537)	8788-6792(	NDC:60432-
Route of Admi	nistration	ORAL					
Active Ingree	dient/Active	Moiety					
	Ingredient	Name	Basi	s of Stren	gth	Stre	ngth
Nystatin (UNII: BI	DF101C72E) (Nys	tatin - UNII:BDF101C72E)	Nystati	n	10	00000 [USP	''U] in 1 mL
Inactive Ingr	edients						
		Ingredient Name				S	trength
water (UNII: 0590	F0KO0R)						
		UNII: 6M3P64V0NC)					
propylene glyco		7V3)					
glycerin (UNII: PD							
sucrose (UNII: C1							
		<b>E</b> (UNII: CI71S98N1Z)					
alcohol (UNII: 3Kg							
methylparaben							
propylparaben ( D&C yellow no.	<b>10</b> (UNII: 355W51	JSQ3G)					
D&C yellow no.	<b>10</b> (UNII: 355W51	JSQ3G)					
D&C yellow no. FD&C red no. 40	10 (UNII: 355W5U ) (UNII: WZB9127	JSQ3G)					
D&C yellow no. FD&C red no. 40 Product Cha	10 (UNII: 355W50 0 (UNII: WZB9127	JSQ3G) XOA)		Scor	~ <u>~</u>		
D&C yellow no. FD&C red no. 40 Product Cha Color	10 (UNII: 355W50 0 (UNII: WZB9127	JSQ3G)		Scor	-		
D&C yellow no. FD&C red no. 40 <b>Product Cha</b> Color Shape	10 (UNII: 355W50 0 (UNII: WZ B9127 racteristics YELLOW (Lig	JSQ3G) XOA)		Size		do	
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D&C yellow no. FD&C red no. 40 Product Cha Color Shape Flavor Contains Packaging # Item Code 1 NDC:68788-	10 (UNII: 35SW50 ) (UNII: WZ B9127 racteristics YELLOW (Lig FRUIT Pa 60 mL in 1 BOT	JSQ3G) XOA) ht creamy yellow) <b>Ackage Description</b> TLE, PLASTIC; Type 0: Not a		Size Impr Marketin Dat	rint Coo	rt Mark	
D&C yellow no. FD&C red no. 40 Product Char Color Shape Flavor Contains Packaging # Item Code 1 NDC:68788- 6792-6	10 (UNII: 35SW50 UNII: WZ B9127 racteristics YELLOW (Lig FRUIT 60 mL in 1 BOT Combination Pro	JSQ3G) XOA) ht creamy yellow) ackage Description TLE, PLASTIC; Type 0: Not a oduct		Size Impr Marketin Dat	rint Coo	rt Mark	
D&C yellow no. FD&C red no. 40 Product Cha Color Shape Flavor Contains Packaging # Item Code 1 NDC:68788-	10 (UNII: 35SW50 (UNII: WZ B9127 racteristics YELLOW (Lig FRUIT 60 mL in 1 BOT Combination Pro	JSQ3G) XOA) ht creamy yellow) ackage Description TLE, PLASTIC; Type 0: Not a oduct		Size Impr Marketin Dat	ng Start	rt Mark	eting End Date

Labeler - Preferred Pharmaceuticals, Inc. (791119022)

**Registrant -** Preferred Pharmaceuticals, Inc. (791119022)

Establishment									
Name	Address	ID/FEI	<b>Business Operations</b>						
Preferred Pharmaceuticals, Inc.		791119022	RELABEL(68788-6792)						

Revised: 7/2024

Preferred Pharmaceuticals, Inc.