

**NYSTATIN- nystatin suspension**  
**ATLANTIC BIOLOGICALS CORP.**

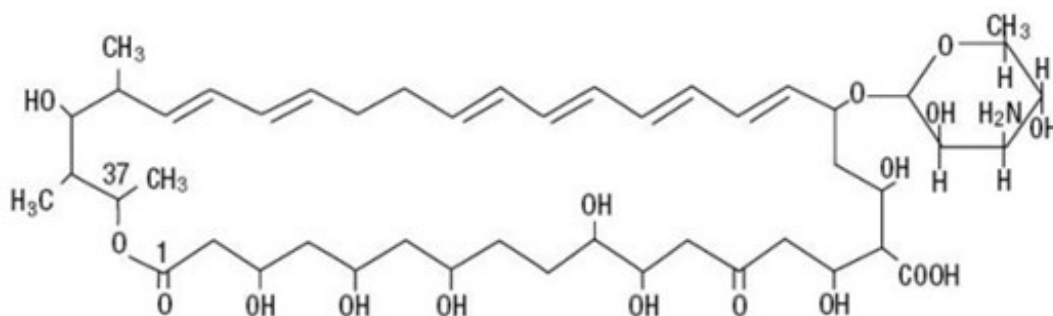
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**NYSTATIN ORAL**  
**SUSPENSION, USP**

**(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 with a calibrated dropper and 1 Pint (473 mL) bottles.

### **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**

28537B  
REV. 09-09

## **PRINCIPAL DISPLAY PANEL - 60 mL Bottle Carton**

### **MGP**

NDC 17856-0037-1

**NYSTATIN ORAL  
SUSPENSION, USP  
(100,000 units  
per mL)**

**Fruit Flavored**

**SHAKE WELL BEFORE USING**

**Packaged with Calibrated Dropper**

**SAFETY SEALED BOTTLE**

**Rx Only**

NET: 2 fl oz (60 mL)

**NDC 17856-0037-01**

NYSTATIN ORAL  
SUSPENSION, USP  
100,000 units per mL

**Rx Only**  
FRUIT FLAVORED  
UNIT DOSE 5 mL Cup

**PACKAGING INFORMATION:**

Dosage per Cup: 5 mL  
Cup(s) per Case: 72

See package insert for indications and dosage schedule.

*Other information:*

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

AVOID FREEZING

**SHAKE WELL BEFORE USING**

**KEEP NYSTATIN ORAL SUSPENSION AND ALL  
DRUGS OUT OF REACH OF CHILDREN**

Mfg by: Morton Grove Pharmaceuticals, Inc.  
Morton Grove, IL 60053  
Repackaged by: UDose LLC, Miami, FL 33179  
Distributed by: Atlantic Biologicals Corp.  
20101 N.E. 16th Place  
Miami, FL 33179

\*Retain box label and package insert for drug information.

Questions or Comments:  
Call 1-800-509-7592

UDose Lot No: XXXXXX  
Mfg Lot No: XXXXXX  
Exp. Date: XX/XX/XXXX



17856003701

## NYSTATIN

nystatin suspension

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:17856-0037(NDC:60432-537)
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Nystatin (UNII: BDF1O1C72E) (Nystatin - UNII:BDF1O1C72E)	Nystatin	100000 [USP <sup>U</sup> ] in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0K00R)	
magnesium aluminum silicate (UNII: 6M3P64V0NC)	
propylene glycol (UNII: 6DC9Q167V3)	
glycerin (UNII: PDC6A3C0OX)	
sucrose (UNII: C151H8M554)	
potassium phosphate, dibasic (UNII: CI71S98N1Z)	

<b>alcohol</b> (UNII: 3K9958V90M)	
<b>methylparaben</b> (UNII: A2I8C7HI9T)	
<b>propylparaben</b> (UNII: Z8IX2SC1OH)	
<b>D&amp;C yellow no. 10</b> (UNII: 35SW5USQ3G)	
<b>FD&amp;C red no. 40</b> (UNII: WZB9127XOA)	

### Product Characteristics

<b>Color</b>	YELLOW (Light creamy yellow)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	FRUIT	<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17856-0037-1	5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product	09/04/2016	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA062512	04/15/1995	

**Labeler** - ATLANTIC BIOLOGICALS CORP. (047437707)

### Establishment

Name	Address	ID/FEI	Business Operations
ATLANTIC BIOLOGICALS CORP.		047437707	repack(17856-0037)

Revised: 9/2016

ATLANTIC BIOLOGICALS CORP.