

**NYSTATIN- nystatin tablet, coated**  
**Heritage Pharmaceuticals Inc**

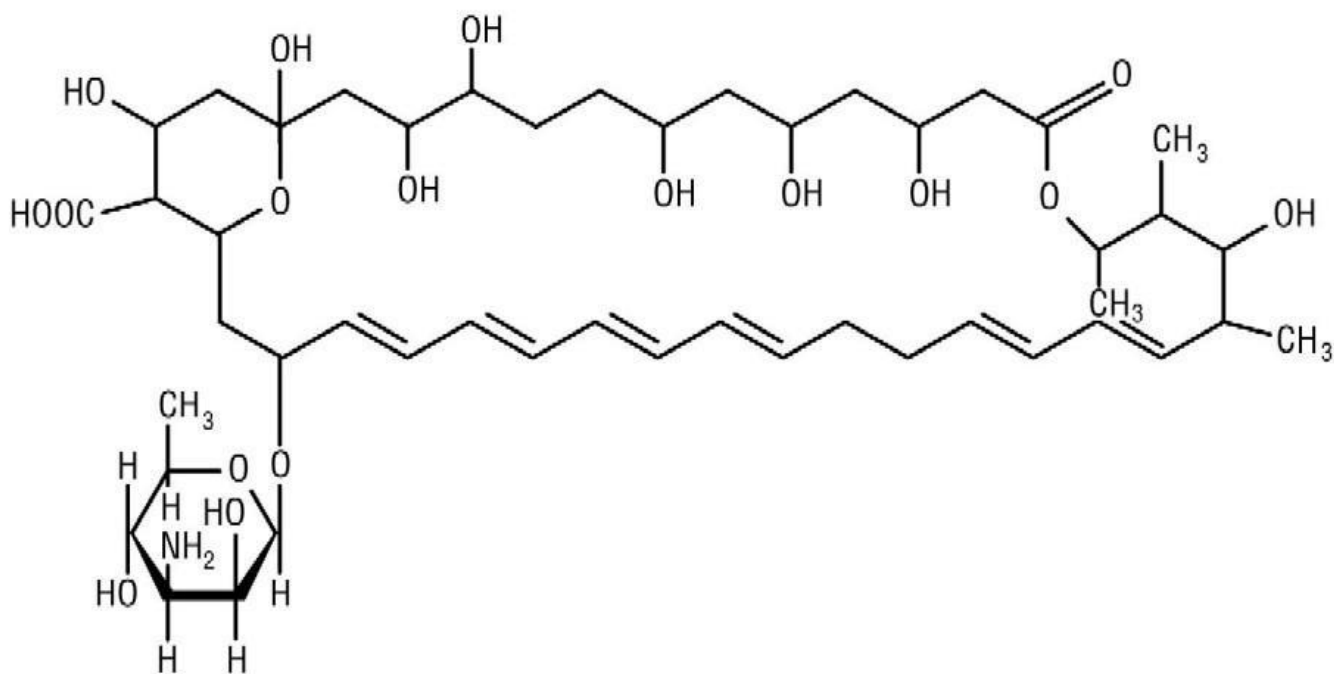
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Nystatin-3/23/2016

**These highlights do not include all the information needed to use Nystatin safely and effectively. see full prescribing information for Nystatin**

**Nystatin (Nystatin) TABLET, COATED for ORAL use.**  
**Initial U.S. Approval:**

**DESCRIPTION**

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



**CHNO**

**M.W. 926.13**

Nystatin Tablets are for oral administration and contain 500,000 units of nystatin per tablet.

Nystatin Tablets contain the inactive ingredients: corn starch, confectioner sugar, hydroxypropyl cellulose, dibasic calcium phosphate, microcrystalline cellulose, talc, magnesium stearate, hypromellose, polyethylene glycol, titanium dioxide, FD&C yellow #6, FD&C red #40, FD&C blue # 2 and polysorbate 80.

**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 Tablets are intended for the treatment of non-esophageal mucus membrane gastrointestinal candidiasis.

## **CONTRAINDICATIONS**

Nystatin Tablets are contraindicated in patients with a history of hypersensitivity to any of their 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. It is also not known whether nystatin can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. Nystatin 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.

## **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 nonspecific 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**

The usual therapeutic dosage is one to two tablets (500,000 to 1,000,000 units nystatin) three times daily. Treatment should generally be continued for at least 48 hours after clinical cure to prevent relapse.

### **HOW SUPPLIED**

Nystatin Tablets USP, 500,000 Units are brown, film-coated tablets debossed "HP51" on one side and plain on the other side are packaged in:

bottles of 100: NDC 23155-051-01

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

Manufactured for:

**Heritage Pharmaceuticals Inc.**

Eatontown, NJ 07724

1.866.901.DRUG (3784)

Made in India.

PON/DRUGS/16 13 4193

020003330 Revised 02/16

### **PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**

**Nystatin Tablets USP, 500,000 Units, 100 count bottles**

NDC 23155-051-01

# Nystatin Tablets, USP (Oral)

**500,000 Units**

100 Tablets

Rx only



**EACH TABLET CONTAINS:**

Nystatin USP.....500,000 Units.

**Usual Adult Dosage:**

Read Accompanying Literature.

**STORE AT 20° TO 25°C (68° TO 77°F). [See USP Controlled Room Temperature].**

Dispense in a tight, light resistant container as defined in the USP.

Manufactured for:

Heritage Pharmaceuticals Inc.

Easton, NJ 07724

1.886.901.DRUG (3784)

Made in INDIA

PO#/DRUGS/16 13 4193

020003331

02/16

Non Varnish Area



## NYSTATIN

nystatin tablet, coated

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:23155-051
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NYSTATIN (UNII: BDF1O1C72E) (NYSTATIN - UNII:BDF1O1C72E)	NYSTATIN	500000 [USP'U]

### Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
SUCROSE (UNII: C151H8M554)	
HYDROXYPROPYL CELLULOSE (UNII: RFW2ET671P)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ANHYDROUS DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
TALC (UNII: 7SEV7J4R1U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
HYPROMELLOSE 2910 (6 MP.A.S) (UNII: 0WZ8WG20P6)	
HYPROMELLOSE 2910 (3 MP.A.S) (UNII: 0VUT3PMY82)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYETHYLENE GLYCOL 8000 (UNII: Q662QK8M3B)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	

**Product Characteristics**

<b>Color</b>	BROWN	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	10mm
<b>Flavor</b>		<b>Imprint Code</b>	HP;51
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:23155-051-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	10/31/2011	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA062474	10/31/2011	

**Labeler** - Heritage Pharmaceuticals Inc (780779901)**Establishment**

Name	Address	ID/FEI	Business Operations
Strides Shasun Ltd.		915786829	MANUFACTURE(23155-051)

Revised: 3/2016

Heritage Pharmaceuticals Inc