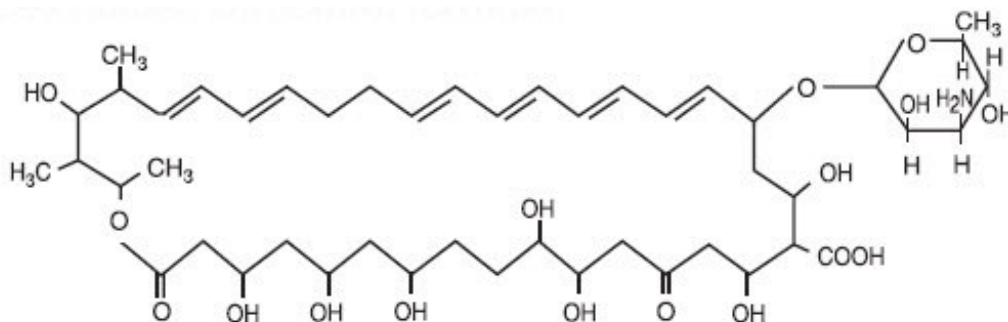

Rx only

DESCRIPTION:

Molecular Formula C₄₇H₇₅NO₁₇

Nystatin cream for topical use, contains 100,000 USP nystatin units per gram in an aqueous, vanishing cream base containing Polysorbate 60, Aluminum Hydroxide Compressed Gel, Titanium Dioxide, Glyceryl Monostearate, Polyethylene Glycol 400 Monostearate, Simethicone Emulsion, Sorbic Acid, Propylene Glycol, Ethylenediamine, Polyoxyethylene Fatty Alcohol Ether, Sorbitol Solution, Methylparaben, Propylparaben, Hydrochloric Acid, White Petrolatum, and Purified Water.

Pharmacokinetics: Nystatin is not absorbed from intact skin or mucous membrane.

INDICATIONS AND USAGE:

Nystatin cream is indicated in the treatment of cutaneous or mucocutaneous mycotic infections caused by *Candida albicans* and other susceptible *Candida* species.

This cream is not indicated for systemic, oral, intravaginal or ophthalmic use.

CONTRAINDICATIONS:

Nystatin cream is contraindicated in patients with a history of hypersensitivity to any of their components.

PRECAUTIONS:

General: Nystatin cream should not be used for the treatment of systemic, oral, intravaginal or ophthalmic infections.

If irritation or sensitization develops, treatment should be discontinued and appropriate measures taken as indicated. It is recommended that KOH smears, cultures, or other diagnostic methods be used to confirm the diagnosis of cutaneous or mucocutaneous candidiasis and to rule out infection caused by other pathogens.

INFORMATION FOR THE PATIENT: Patients using this medication should receive the following information and instructions:

1. The patient should be instructed to use this medication as directed (including the replacement of missed doses). This medication is not for any disorder other than that for which it is prescribed.
2. Even if symptomatic relief occurs within the first few days of treatment, the patient should be advised not to interrupt or discontinue therapy until the prescribed course of treatment is completed.
3. If symptoms of irritation develop, the patient should be advised to notify the physician promptly.

Laboratory Tests: If there is a lack of therapeutic response, KOH smears, cultures, or other diagnostic methods should be repeated.

Carcinogenesis, Mutagenesis, Impairment of Fertility: No long-term animal studies have been performed to evaluate the carcinogenic potential of nystatin. No studies have been performed to determine the mutagenicity of nystatin or the effects on male or female fertility.

Pregnancy: Teratogenic Effects: Category C: Animal reproduction studies have not been conducted with any nystatin cream. It also is not known whether this cream can cause fetal harm when used by a pregnant woman or can affect reproductive capacity. Nystatin cream should be prescribed for a pregnant woman only if the potential benefit to the mother outweighs the potential risk to the fetus.

Nursing Mothers: It is not known whether nystatin is excreted in human milk. Caution should be exercised when nystatin is prescribed for a nursing woman.

Pediatric Use: Safety and effectiveness have been established in the pediatric population from birth to 16 years. (See DOSAGE AND ADMINISTRATION.)

Geriatric Use: Clinical studies with nystatin cream did not include sufficient numbers of subjects aged 65 years and older to determine whether they respond differently than younger subjects. Other reported clinical experience has not identified differences in responses between elderly and younger patients, but greater sensitivity of some older individuals cannot be ruled out.]

ADVERSE REACTIONS:

The frequency of adverse events reported in patients using nystatin cream is less than 0.1%. The more common events that were reported include allergic reactions, burning, itching, rash, eczema, and pain on application. (See PRECAUTIONS: General.)

DOSAGE AND ADMINISTRATION:

Adults and Pediatric Patients (Neonates and Older): Apply liberally to affected area twice daily or as indicated until healing is complete.

HOW SUPPLIED:

Nystatin Cream USP is available as follows:

NDC 0168-0054-15 15 gram tube

NDC 0168-0054-30 30 gram tube

Each tube provides 100,000 USP Nystatin Units per gram.

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

Avoid freezing.

E. FOUGERA & CO.

A division of

Fougera

PHARMACEUTICALS INC.

Melville, New York 11747

I25415G

R07/13

#61

PRINCIPAL DISPLAY PANEL - 15 GM Tube Carton Label

CommUnity Care Federally Qualified Centers

NYSTATIN

CREAM

15 GM

Date:

Name:

Dr.

APPLY TO AFFECTED AREA 2 TIMES DAILY.

123456

1/1/01

NYSTATIN CRM 15 gm NDC 76413-119-15

Batch: 123456

Lot: 123456

Exp: 1/1/01

FOUGERA

Federal law prohibits the transfer of this drug to any other person than the patient for whom prescribed.

CommUnity Care Federally Qualified Centers

NYSTATIN
CREAM
15 GM

Date:

Name: Dr.

APPLY TO AFFECTED AREA 2 TIMES DAILY.

APLIQUE AL AREA AFECTADA 2 VECES AL DIA.

123456

1/1/01

NYSTATIN CRM 15 gm NDC 76413-119-15

Batch: 123456 Lot: 123456 Exp: 1/1/01 FOUGERA
Federal law prohibits the transfer of this drug to any other person than the patient for whom prescribed.

NYSTATIN
nystatin cream

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:76413-119(NDC:0168-0054)
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
nystatin (UNII: BDF1O1C72E) (nystatin - UNII:BDF1O1C72E)		nystatin	100000 [USP.U] in 1 g
Inactive Ingredients			
Ingredient Name			Strength
Polysorbate 60 (UNII: CAL22UVI4M)			
Algeldrate (UNII: 03J11K103C)			
Titanium Dioxide (UNII: 15FIX9V2JP)			
Glyceryl Monostearate (UNII: 230OU9XXE4)			
PEG-8 Stearate (UNII: 2P9L47VI5E)			
Silicon Dioxide (UNII: ETJ7Z6XBU4)			
Sorbic Acid (UNII: X045WJ989B)			
Propylene Glycol (UNII: 6DC9Q167V3)			
Ethylenediamine (UNII: 60V9STC53F)			
Sorbitol (UNII: 506T60A25R)			
Methylparaben (UNII: A2I8C7HI9T)			
Propylparaben (UNII: Z8IX2SC1OH)			
Hydrochloric Acid (UNII: QTT17582CB)			

Petrolatum (UNII: 4T6H12BN9U)				
Water (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76413-119-15	1 in 1 CARTON		
1		15 g in 1 TUBE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA062129		02/01/1979	

Labeler - Central Texas Community Health Centers (079674019)

Establishment				
Name	Address	ID/FEI	Business Operations	
Central Texas Community Health Centers		079674019	REPACK(76413-119) , RELABEL(76413-119)	