CANDIDA ALBICANS PHENOLIC- candida albicans liquid Energique, Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

DRUG FACTS:

ACTIVE INGREDIENT:

(in each drop): 25% of Candida Albicans 12X, 30X, 12C, 30C.

INDICATIONS:

May temporarily relieve symptoms associated with reactions to candida albicans, such as indigestion.**

**Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

WARNINGS:

If pregnant or breastfeeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Do not use if tamper evident seal is broken or missing.

Store in a cool, dry place.

KEEP OUT OF REACH OF CHILDREN:

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Adults and children 5 to 10 drops orally, 1 time daily or as otherwise directed by a health care professional. If symptoms persist for more than 7 days, consult your health care professional. Consult a physician for use in children under 12 years of age.

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FDA evaluated.

INACTIVE INGREDIENTS:

Demineralized water, 20% Ethanol.

QUESITONS:

Dist. by Energique, Inc. 201 Apple Blvd. Woodbine, IA 51579 **800.869.8078**

PACKAGE LABEL DISPLAY:

ENERGIQUE

SINCE 1987

HOMEOPATHIC REMEDY

CANDIDA

ALBICANS

PHENOLIC

1 fl. oz. (30 ml)

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LOT: XXXXXX MFD: MM/YY



CANDIDA ALBICANS PHENOLIC candida albicans liquid							
Product Information							
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0237				
Route of Administration	ORAL						

Active Ingredient/Active Moiety								
		Ingredient Name		Basis of Strength	Strength			
	ANDIDA ALBICA	NS (UNII: 4D7G21HDBC) (CANDIDA ALBICANS -		CANDIDA ALBICANS	5 12 [hp_X] in 1 mL			
In	nactive Ingr	edients						
	Ingredient Name S							
W	ATER (UNII: 059	QF0KO0R)						
AL	COHOL (UNII: 3	K9958V90M)						
P	ackaging							
#	Item Code	Package Description	Ma	rketing Start Date	Marketing End Date			
1	NDC:44911- 0237-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/19	9/2015				
Marketing Information								
	Marketing Category	Application Number or Monograph Citation	Marl	keting Start Date	Marketing End Date			
	approved meopathic		08/19/2	2015				

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment						
Name	Address	ID/FEI	Business Operations			
Apotheca Company		844330915	manufacture(44911-0237) , api manufacture(44911-0237) , label(44911-0237) , pack(44911-0237)			

Revised: 7/2023

Energique, Inc.