CANDIDA- candida albicans liquid Deseret Biologicals, 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:

Candida Albicans 200C.

HOMEOPATHIC INDICATIONS:

For temporary relief of symptoms related to Candida Albicans infection including nausea, drowsiness, lethargy, confusion, vaginal discharge, sensitivities to foods and other fungi, petrochemicals, and mucous congestion.**

**These statements are based upon traditional homeopathic principles. They have not been reviewed by the Food and Drug Administration.

WARNINGS:

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

If pregnant or breast-feeding, ask a health professional before use.

Tamper seal: "Sealed for Your Protection." Do not use if seal is broken or missing.

KEEP OUT OF REACH OF CHILDREN:

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

DIRECTIONS:

1-10 drops under the tongue, 3 times a day or as directed by a health professional. Consult a physician for use in children under 12 years of age.

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INACTIVE INGREDIENTS:

Demineralized Water, 25% Ethanol

QUESTIONS:

Dist. By: Deseret Biologicals, Inc.

469 W. Parkland Drive

Sandy, UT 84070 www.desbio.com

PACKAGE LABEL DISPLAY:

DESBIO

NDC 43742-0985-1

HOMEOPATHIC

CANDIDA 200C

1 FL OZ (30 ml)

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CANDIDA

candida albicans liquid

Product Type HUMAN OTC DRUG Item Code (Source) NDC:43742-0985

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength CANDIDA ALBICANS (UNII: 4D7G21HDBC) (CANDIDA ALBICANS - UNII: 4D7G21HDBC) CANDIDA ALBICANS CANDIDA ALBICANS CANDIDA ALBICANS 200 [hp_C] in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
ALCOHOL (UNII: 3K9958V90M)				

Packaging					
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:43742-0985-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	02/21/2017	10/13/2025	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved homeopathic		02/21/2017	10/13/2025		

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0985), api manufacture(43742-0985), label(43742-0985), pack(43742-0985)

Revised: 7/2021 Deseret Biologicals, Inc.