TOXOPLASMA GONDII HOMOCHORD- toxoplasma gondii 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 INGREDIENTS:

Toxoplasma Gondii 15X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X.

HOMEOPATHIC INDICATIONS:

For the temporary relief of symptoms of body aches, headache, fever, and fatigue **

**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 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 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.

HOMEOPATHIC INDICATIONS:

For the temporary relief of symptoms of body aches, headache, fever, and fatigue **

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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:

DES**BIO**

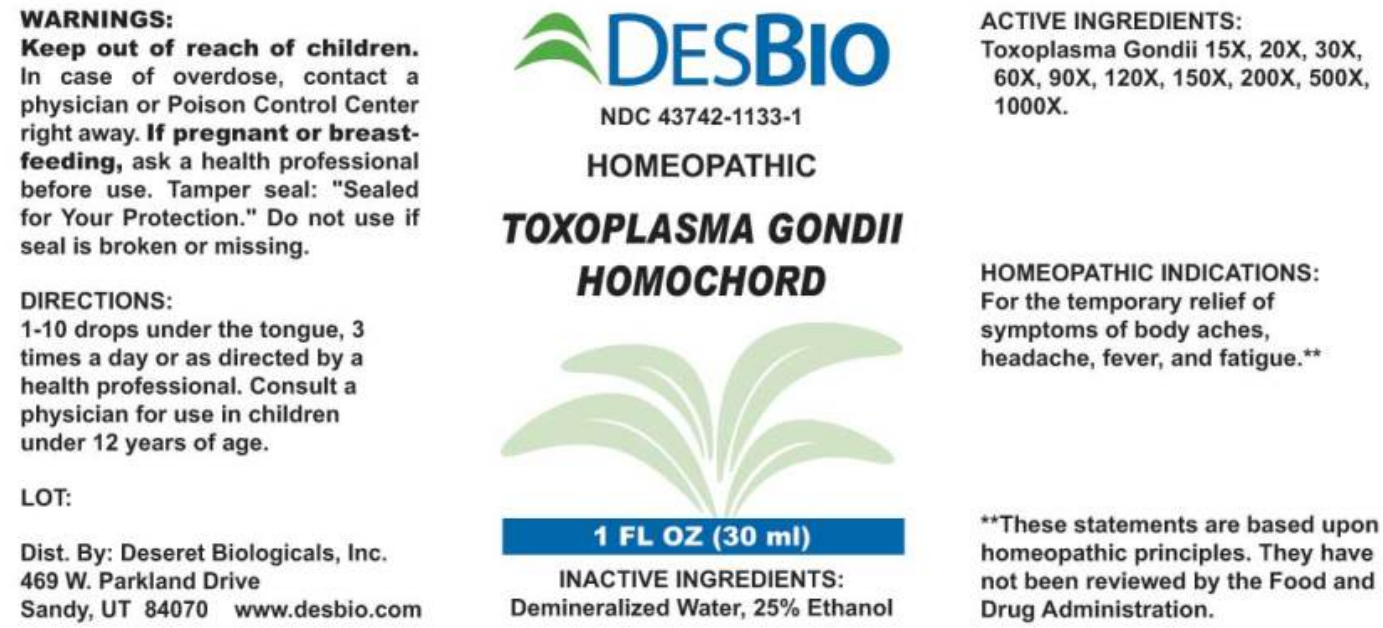
NDC 43742-1133-1

HOMEOPATHIC

TOXOPLASMA GONDII

HOMOCHORD

1 FL OZ (30 ml)



TOXOPLASMA GONDII HOMOCHORD

toxoplasma gondii liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43742-1133
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TOXOPLASMA GONDII (UNII: BMV90JF469) (TOXOPLASMA GONDII - UNII:BMV90JF469)	TOXOPLASMA GONDII	15 [hp_X] 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:43742-1133-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	12/29/2017	02/28/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/29/2017	02/28/2025

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-1133) , api manufacture(43742-1133) , label(43742-1133) , pack(43742-1133)

Revised: 8/2021 Deseret Biologicals, Inc.