

**TOXO HOMO- toxoplasma gondii dna 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.*

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**Drug Facts:**

**ACTIVE INGREDIENTS:**

Toxoplasma Gondii DNA 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 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.

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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 43742-2002-1**

**HOMEOPATHIC**

**TOXO:HOMO**

**1 FL OZ (30 ml)**

**WARNINGS:**

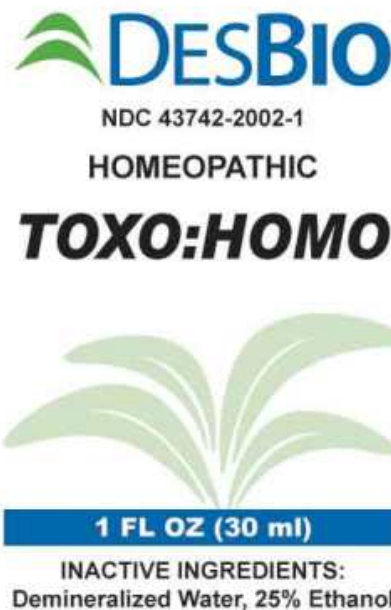
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**LOT:**

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**TOXO HOMO**

toxoplasma gondii dna liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:43742-2002
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
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<b>TOXOPLASMA GONDII DNA</b> (UNII: JMX3ACIU25) (TOXOPLASMA GONDII DNA - UNII:JMX3ACIU25)	TOXOPLASMA GONDII DNA	15 [hp_X] in 1 mL
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### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43742-2002-1	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/21/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		12/21/2020	

**Labeler** - Deseret Biologicals, Inc. (940741853)

**Registrant** - Apotheca Company (844330915)

### Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-2002) , api manufacture(43742-2002) , label(43742-2002) , pack(43742-2002)

Revised: 1/2023

Deseret Biologicals, Inc.