

CHLA HOMO- chlamydia trachomatis 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:

Chlamydia Trachomatis 12X, 15X, 20X, 30X, 60X, 120X, 150X, 200X, 500X, 1000X.

HOMEOPATHIC INDICATIONS:

For temporary relief of symptoms related to chlamydia infections. Male: burning sensation during urination, discharge from penis and testicular tenderness or pain. Female: vaginal discharge, burning sensation during urination, and painful sexual intercourse.**

**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 a 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 a 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 43742-1771-1

HOMEOPATHIC

CHLA:HOMO

HOMOCHORD

1 FL OZ (30 ml)

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LOT:

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CHLA HOMO

chlamydia trachomatis liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-1771
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLAMYDIA TRACHOMATIS (UNII: T6NI39QU44) (CHLAMYDIA TRACHOMATIS - UNII:T6NI39QU44)	CHLAMYDIA TRACHOMATIS	12 [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:43742-1771-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	02/26/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		02/26/2021	

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-1771) , api manufacture(43742-1771) , label(43742-1771) , pack(43742-1771)

Revised: 1/2023

Deseret Biologicals, Inc.