CHLAMYDIA REMEDY- 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.

INDICATIONS:

For temporary relief of symptoms related to chlamydia infections. Male: burning sensation during urination, discharge from penis, testicular tenderness or pain and rectal discharge or pain. Female: vaginal discharge, burning sensation during urination, painful sexual intercourse, rectal pain or discharge, liver pain, dark colored urine and clay colored feces.

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

PACKING LABEL DISPLAY:

DES**BIO**

NDC 43742-0500-1

HOMEOPATHIC

CHLAMYDIA REMEDY

1 FL OZ (30 ml)

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LOT:

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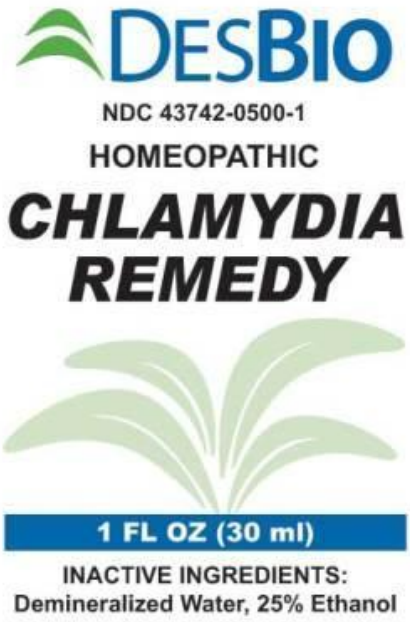

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CHLAMYDIA REMEDY

chlamydia trachomatis liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43742-0500
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:43742-0500-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/10/2014	09/28/2020

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/10/2014	09/28/2020

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0500) , api manufacture(43742-0500) , label(43742-0500) , pack(43742-0500)

Revised: 4/2018 Deseret Biologicals, Inc.