MYCOPLASMA HOMOCHORD- mycoplasma pneumoniae 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:

Mycoplasma Pneumoniae 15X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X.

HOMEOPATHIC INDICATIONS:

For temporary relief of symptoms related to Mycoplasma Infection including headache, muscle and joint pain and soreness, and nausea.**

**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. 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:

QUESTIONS:

Dist. By:

Deseret Biologicals, Inc.

469 W. Parkland Drive

Sandy, UT 84070

www.desbio.com

PACKAGE LABEL DISPLAY:

DESBIO

NDC 43742-0843-1

HOMEOPATHIC

MYCOPLASMA

HOMOCHORD

1 FL OZ (30 ml)

WARNINGS:

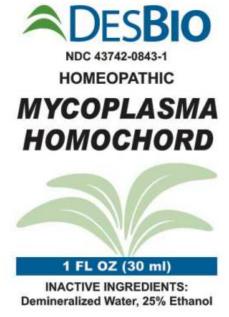
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MYCOPLASMA HOMOCHORD

mycoplasma pneumoniae liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-0843
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MYCOPLASMA PNEUMONIAE (UNII: JQE470FAD0) (MYCOPLASMA PNEUMONIAE - UNII:JQE470FAD0)	MYCOPLASMA PNEUMONIAE	15 [hp_X] in 1 mL	

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

Packaging					
#	Item Code	tem Code Package Description Marketing Star Date		: Marketing End Date	
	NDC:43742- 0843-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/08/2016	07/07/2025	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		09/08/2016	07/07/2025	

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0843) , api manufacture(43742-0843) , label(43742-0843) , pack(43742-0843)

Revised: 8/2021 Deseret Biologicals, Inc.