

HPYL HOMO- helicobacter pylori 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:

Helicobacter Pylori 15X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X.

HOMEOPATHIC INDICATIONS:

For the temporary relief of symptoms related to H. Pylori infection including upper abdominal pain or discomfort, loss of appetite, belching, nausea, vomiting 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 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:

DES BIO

NDC 43742-1848-1

HOMEOPATHIC

HPYL:HOMO

1 FL OZ (30 ml)

WARNINGS:

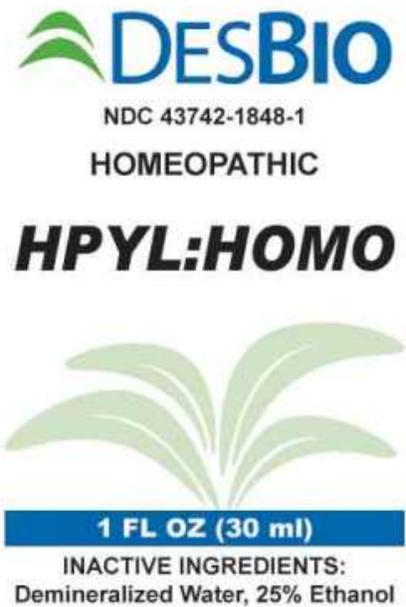
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HPYL HOMO

helicobacter pylori liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-1848
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of	Strength
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Ingredient Name		Strength	Strength	
HELICOBACTER PYLORI (UNII: U09W5JOL3Z) (HELICOBACTER PYLORI - UNII:U09W5JOL3Z)		HELICOBACTER PYLORI	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:43742-1848-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	10/26/2020	11/06/2028
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
unapproved homeopathic			10/26/2020	11/06/2028

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-1848) , api manufacture(43742-1848) , label(43742-1848) , pack(43742-1848)

Revised: 2/2024

Deseret Biologicals, Inc.