# PEPT HOMO- peptostreptococcus anaerobius 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:**

Peptostreptococcus Anaerobius DNA 19X, 20X, 30X, 60X, 90X, 120X, 150X, 200X, 500X, 1000X.

# **HOMEOPATHIC INDICATIONS:**

For temporary relief of symptoms related to Peptostrep infection including nasal or sinus congestion, and earache.\*\*

\*\*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 unde 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 43742-1967-1

**HOMEOPATHIC** 

**PEPT:HOMO** 

1 FL OZ (30 ml)

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#### LOT:

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# **PEPT HOMO**

peptostreptococcus anaerobius dna liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-1967
Route of Administration	ORAL		

# **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
PEPTOSTREPTOCOCCUS ANAEROBIUS DNA (UNII: RKA0GWXVH6) (PEPTOSTREPTOCOCCUS ANAEROBIUS DNA - UNII: RKA0GWXVH6)	PEPTOSTREPTOCOCCUS ANAEROBIUS DNA	19 [hp_X] in 1 mL

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

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:43742- 1967-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	04/13/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		04/13/2021	

# Labeler - Deseret Biologicals, Inc. (940741853)

# Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-1967) , api manufacture(43742-1967) , label(43742-1967) , pack(43742-1967)

Revised: 1/2023 Deseret Biologicals, Inc.