ACETYLCHOLINE CHLORIDE- acetylcholine chloride 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:

Acetylcholine Chloride 6X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

PURPOSE:

For temporary relief of symptoms including dizziness, chest congestion, cough, fatigue, rash, itching, headache and poor memory.**

**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:

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.

INDICATIONS:

For temporary relief of symptoms including dizziness, chest congestion, cough, fatigue, rash, itching, headache and poor memory.**

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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 43742-0817-1

HOMEOPATHIC

ACETYLCHOLINE CHLORIDE

1 FL OZ (30 ml)

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ACETYLCHOLINE CHLORIDE
acetylcholine chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43742-0817
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

ACETYLCHOLINE CHLORIDE (UNII: AF73293C2R) (ACETYLCHOLINE - UNII:N9YNS0M02X)	ACETYLCHOLINE	6 [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:43742-0817-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/21/2016	09/15/2028

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/21/2016	09/15/2028

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0817) , api manufacture(43742-0817) , label(43742-0817) , pack(43742-0817)

Revised: 3/2024 Deseret Biologicals, Inc.