ACETYLCHOLINE CHLORIDE PHENOLIC- acetylcholine chloride, liquid
Energique, 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:

(in each drop): 20% of Acetylcholine Chloride 6X, 12X, 30X, 12C, 30C.

INDICATIONS:

May temporarily relieve symptoms associated with reactions to actylcholine chloride, such as chest congestion with mild constriction. **

**Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

WARNINGS:

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Do not use if tamper evident seal is broken or missing. Store in a cool, dry place.

DIRECTIONS:

Adults and children 5 to 10 drops orally, 1 time daily or as otherwise directed by a health care professional. If symptoms persist for more than 7 days, consult your health care professional. Consult a physician for use in children under 12 years of age.

INACTIVE INGREDIENTS:

Demineralized water, 20% Ethanol.

KEEP OUT OF REACH OF CHILDREN.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

INDICATIONS:

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such as chest congestion with mild constriction. **

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QUESTIONS:

Dist. by Energique, Inc.

201 Apple Blvd

Woodbine, IA 51579 800.869.8078

PACKAGE LABEL DISPLAY

ENERGIQUE

SINCE 1987

HOMEOPATHIC REMEDY

ACETYLCHOLINE

CHLORIDE

PHENOLIC

1 fl. oz. (30 ml)

ACETYLCHOLINE CHLORIDE PHENOLIC

acetylcholine chloride, liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44911-0025
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:44911-0025-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	11/08/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		11/08/2012	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0025) , api manufacture(44911-0025) , label(44911-0025) , pack(44911-0025)

Revised: 6/2021 Energique, Inc.