ACONITE- aconitum napellus 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 INGREDIENT:

(in each drop): 100% of Aconitum Napellus 200C.

INDICATIONS:

May temporarily relieve sudden onset of illness with intense pain.**

**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.

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.

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.

INDICATIONS:

May temporarily relieve sudden onset of illness with intense pain.**

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

INACTIVE INGREDIENTS:

Demineralized water, 20% Ethanol.

QUESTIONS:

Dist. by Energique, Inc.

201 Apple Blvd

Woodbine, IA 51579 **800-869-8078**

PACKAGE LABEL DISPLAY:

ENERGIQUE

SINCE 1987

HOMEOPATHIC REMEDY

ACONITE 200C

1 fl. oz. (30 ml)

ACONITE

aconitum napellus liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44911-0209
Route of Administration	ORAL		

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

ACONITUM NAPELLUS WHOLE (UNII: U0NQ8555JD) (ACONITUM NAPELLUS - UNII:U0NQ8555JD)	ACONITUM NAPELLUS WHOLE	200 [hp_C] 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:44911-0209-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/15/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/15/2015	

Labeler - ENERGIQUE, INC. (789886132)

Registrant - APOTHECA COMPANY (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
APOTHECA COMPANY		844330915	manufacture(44911-0209) , api manufacture(44911-0209) , label(44911-0209) , pack(44911-0209)

Revised: 6/2023 ENERGIQUE, INC.