

GABA PHENOLIC- gamma-aminobutyric acid 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 Gamma-Aminobutyric Acid 6X, 12X, 30X, 12C, 30C.

INDICATIONS:

May temporarily relieve symptoms associated with reactions to GABA.**

**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.

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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

GABA PHENOLIC

1 fl. oz. (30 ml)

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LOT: XXXXXX MFD: MM/YY



GABA PHENOLIC

gamma-aminobutyric acid liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0002
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
.GAMMA.-AMINOBTYRIC ACID (UNII: 2ACZ6IPC6I) (.GAMMA.-AMINOBTYRIC ACID - UNII:2ACZ6IPC6I)		.GAMMA.-AMINOBTYRIC ACID	6 [hp_X] in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
ALCOHOL (UNII: 3K9958V90M)				
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44911-0002-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	04/27/2012	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		04/27/2012		

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0002) , api manufacture(44911-0002) , label(44911-0002) , pack(44911-0002)

Revised: 3/2023

Energique, Inc.