NOREPINEPHRINE PHENOLIC- norepinephrine (bitartrate) 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): 25% of Norepinephrine (Bitartrate) 6X, 12X, 30X; 12.50% of Norepinephrine (Bitartrate) 12C, 30C.

INDICATIONS:

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

**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:

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 symptoms associated with reactions to norepinephrine.**

**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

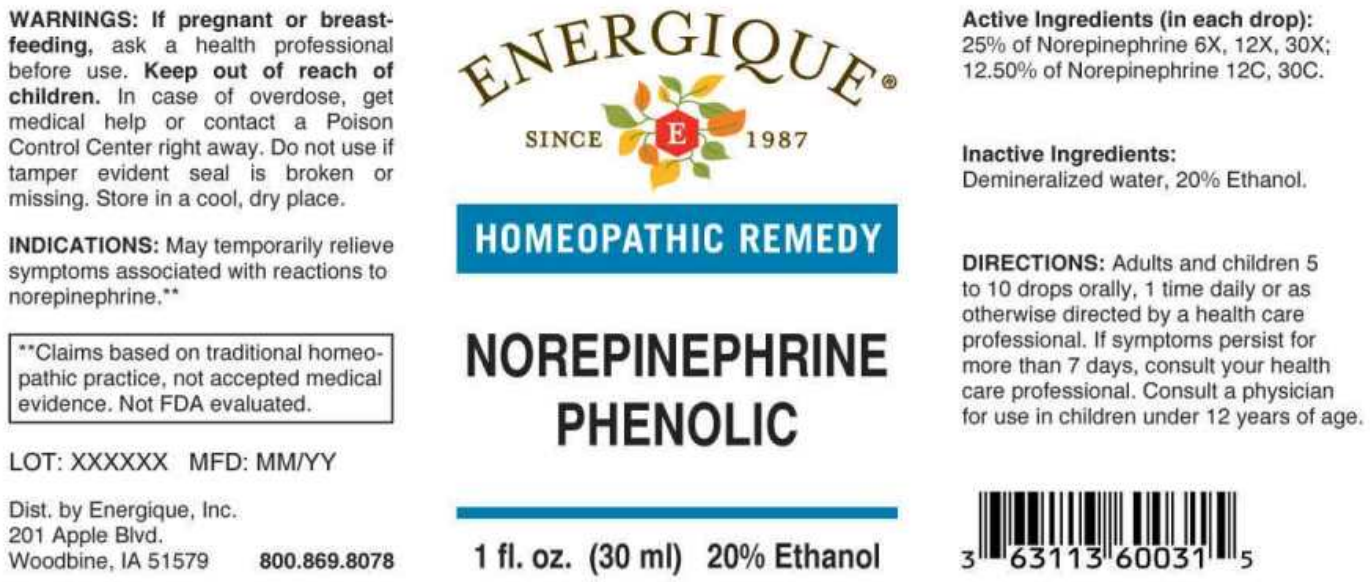
SINCE 1987

HOMEOPATHIC REMEDY

NOREPINEPHRINE

PHENOLIC

1 fl. oz. (30 ml)



NOREPINEPHRINE PHENOLIC

norepinephrine (bitartrate) liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44911-0499
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NOREPINEPHRINE BITARTRATE (UNII: IFY5PE3ZRW) (NOREPINEPHRINE - UNII:X4W3ENH1CV)	NOREPINEPHRINE	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:44911-0499-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	06/07/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		06/07/2019	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0499) , api manufacture(44911-0499) , label(44911-0499) , pack(44911-0499)

Revised: 3/2024 Energique, Inc.