NOREPINEPHRINE PHENOLIC- norepinephrine 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:

Norepinephrine 6X, 12X, 30X, 12C, 30C.

INDICATIONS:

To be used according to standard homeopathic indications.**

**These statements are based upon traditional homeopathic practice. They have not been reviewed by the Food and Drug Administration.

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, consult your health care professional.

Consult a physician for use in children under 12 years of age.

INDICATIONS:

To be used according to standard homeopathic indications.**

**These statements are based upon traditional homeopathic practice. They have not been reviewed by the Food and Drug Administration.

INACTIVE INGREDIENTS:

Demineralized water, 20% Ethanol.

QUESTIONS:

Dist. by Energique, Inc.

201 Apple Blvd

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

PACKAGE LABEL DISPLAY:

ENERGIQUE

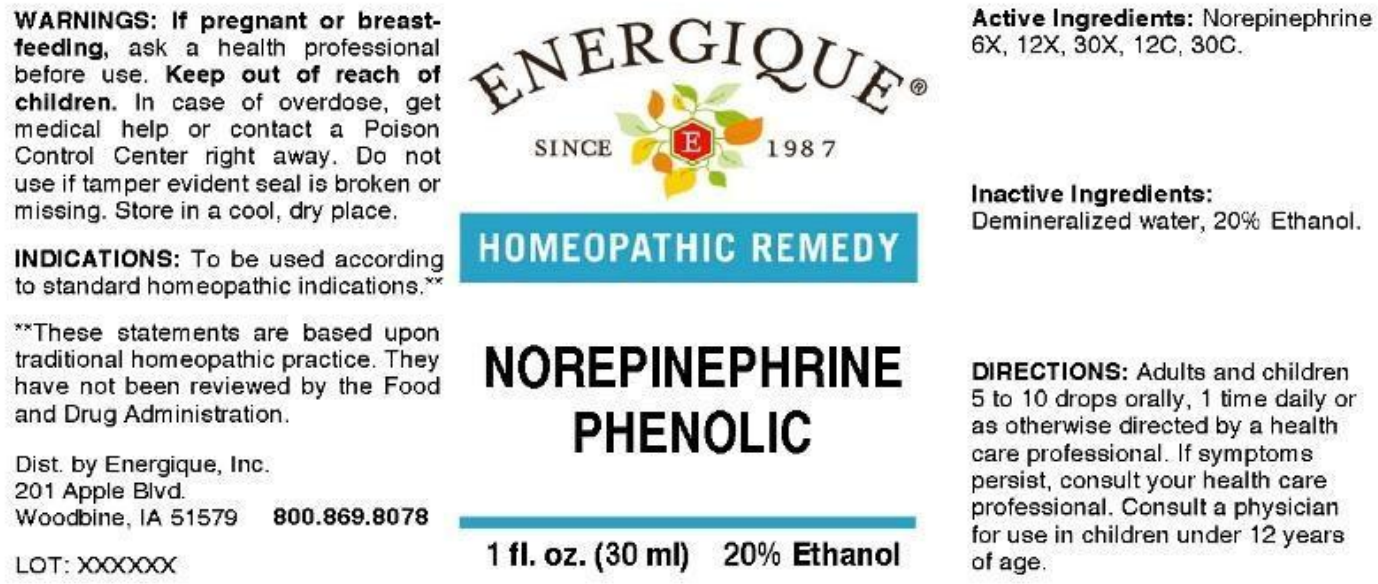
SINCE 1987

HOMEOPATHIC REMEDY

NOREPINEPHRINE

PHENOLIC

1 fl. oz. (30 ml)



NOREPINEPHRINE PHENOLIC

norepinephrine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44911-0282
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NOREPINEPHRINE (UNII: X4W3ENH1CV) (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:44911-0282-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	10/26/2015	05/16/2022

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/26/2015	05/16/2022

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0282) , api manufacture(44911-0282) , label(44911-0282) , pack(44911-0282)

Revised: 4/2020 Energique, Inc.