QUERCETIN PHENOLIC- quercetin, 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): 33.33% of Quercetin 6X; 16.67% of Quercetin 12X, 30X, 12C, 30C.

INDICATIONS:

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

**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:

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

INDICATIONS:

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

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

QUESTIONS:

Dist. by Energique, Inc.

201 Apple Blvd.

Woodbine, IA 51579 800.869.8078

PACKAGE LABEL DISPLAY:

ENERGIQUE

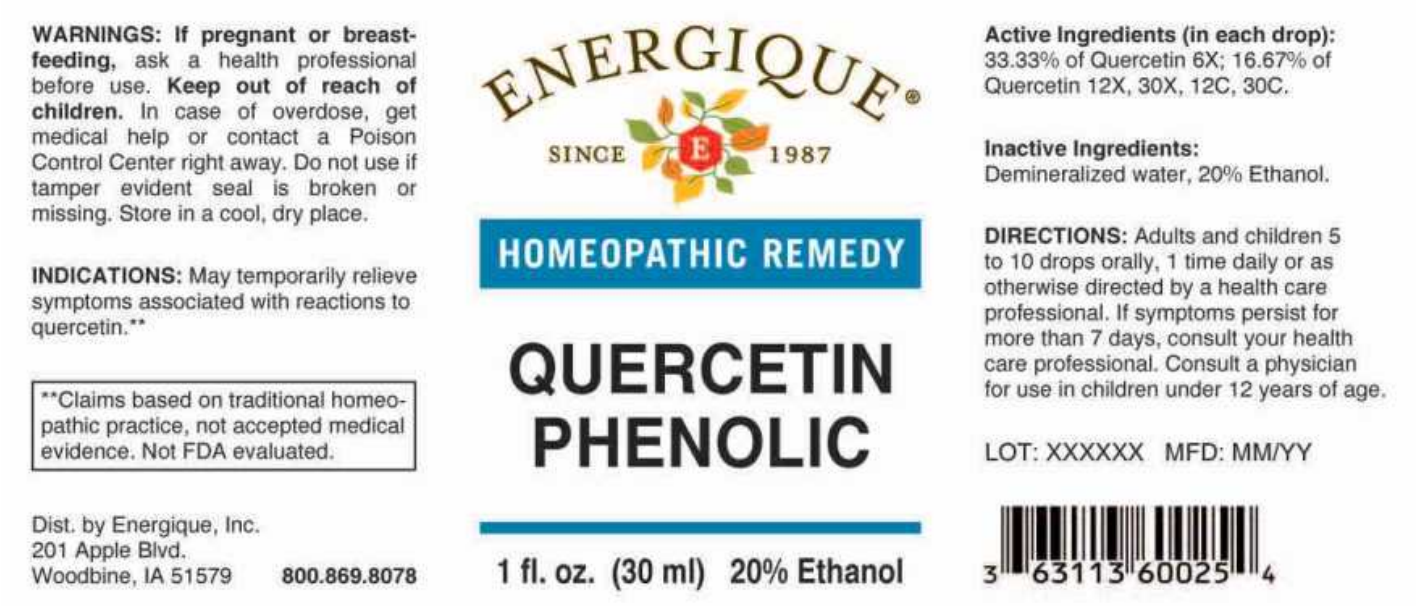
SINCE 1987

HOMEOPATHIC REMEDY

QUERCETIN

PHENOLIC

1 fl. oz. (30 ml)



QUERCETIN PHENOLIC

quercetin, liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44911-0013
Route of Administration	ORAL		

Active Ingredient/Active Moiety		

Ingredient Name	Basis of Strength	Strength
QUERCETIN (UNII: 9IKM0I5T1E) (QUERCETIN - UNII:9IKM0I5T1E)	QUERCETIN	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:44911-0013-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	06/05/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		06/05/2012	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0013) , api manufacture(44911-0013) , label(44911-0013) , pack(44911-0013)

Revised: 3/2024 Energique, Inc.