MENADIONE PHENOLIC- menadione 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): 24.98% of Menadione 12X, 30X, 12C, 30C. 0.10% of Menadione 6X.

INDICATIONS:

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

**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.

Woodbine, IA 51579 800-869-8078

PACKAGE LABEL DISPLAY:

ENERGIQUE

SINCE 1987

HOMEOPATHIC REMEDY

MENADIONE

PHENOLIC

1 fl. oz. (30 ml)

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LOT: XXXXXXX

MENADIONE PHENOLIC

menadione liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:449 11-0 550

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENADIO NE (UNII: 723JX6CXY5) (MENADIO NE - UNII:723JX6CXY5)	MENADIONE	6 [hp_X] in 1 mL

Inactive Ingredients

ı	Ingredient Name	Strength

WATER (UNII: 059QF0KO0R)

ALCOHOL (UNII: 3K9958V90M)

Packaging				
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44911- 0550-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	06/11/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		06/11/2020	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0550), api manufacture(44911-0550), label(44911-0550), pack(44911-0550)

Revised: 6/2020 Energique, Inc.