

**ESTROGEN PHENOLIC- estradiol, 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.*

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**DRUG FACTS:**

**ACTIVE INGREDIENT:**

**(in each drop):** 24.98% of Estradiol 12X, 30X, 12C, 30C; 0.10% of Estradiol 6X.

**INDICATIONS:**

May temporarily relieve symptoms associated with reactions to estrogen, such as breast tenderness and menopausal symptoms.\*\*

\*\*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**

**ESTROGEN**

**PHENOLIC**

**1 fl. oz. (30 ml)**

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LOT: XXXXXX MFD: MM/YY



**ESTROGEN PHENOLIC**

estradiol, liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:44911-0199
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ESTRADIOL (UNII: 4TI98Z838E) (ESTRADIOL - UNII:4TI98Z838E)	ESTRADIOL	6 [hp_X] in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
ALCOHOL (UNII: 3K9958V90M)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44911-0199-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/09/2015	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/09/2015	

**Labeler** - Energique, Inc. (789886132)**Registrant** - Apotheca Company (844330915)**Establishment**

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0199) , api manufacture(44911-0199) , label(44911-0199) , pack(44911-0199)

Revised: 10/2024

Energique, Inc.