

**HOUSEHOLD ANTIGENS-** acetone, sodium tripolyphosphate, propylene glycol, benzinum, ammonium muriaticum, sodium lauryl sulfate, terebinthina, butyl acetate, ethyl acetate, toluene, xylene, bisphenol a, petroleum, anacardium orientale, graphites, nitricum acidum, pulsatilla (pratensis), sulphur 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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## **Drugs:**

### **ACTIVE INGREDIENTS:**

**(in each drop):** 9.09% of Anacardium Orientale 12X, 15X, Graphites 12X, 15X, Nitricum Acidum 12X, 15X, Pulsatilla (Pratensis) 12X, 15X, Sulphur 12X, 15X; 2.27% of Acetone 9X, 12X, 15X, 30X, Sodium Tripolyphosphate 9X, 12X, 15X, 30X, Propylene Glycol 9X, 12X, 15X, 30X, Benzinum 9X, 12X, 15X, 30X, Ammonium Muriaticum 9X, 12X, 15X, 30X, Sodium Lauryl Sulfate 9X, 12X, 15X, 30X, Terebinthina 9X, 12X, 15X, 30X, Butyl Acetate 9X, 12X, 15X, 30X, Ethyl Acetate 9X, 12X, 15X, 30X, Toluene 9X, 12X, 15X, 30X, Xylene 9X, 12X, 15X, 30X, Bisphenol A 9X, 12X, 15X, 30X, Petroleum 9X, 12X, 15X, 30X.

### **INDICATIONS:**

May temporarily relieve symptoms associated with household chemicals sensitivities, such as dizziness and headache.\*\*

\*\*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, 3 times 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 household chemicals sensitivities, such as dizziness and headache.\*\*

\*\*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**

**HOUSEHOLD ANTIGENS**

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

**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.

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**HOMEOPATHIC REMEDY**

# HOUSEHOLD ANTIGENS™

1 fl. oz. (30 ml) 20% Ethanol

**Active Ingredients (in each drop):**  
9.09% of Anacardium 12X, 15X, Graphites 12X, 15X, Nitricum Ac 12X, 15X, Pulsatilla 12X, 15X, Sulphur 12X, 15X; 2.27% of Household Allersodes (Acetone, Ammon Mur, Benzinum, Bisphenol A, Butyl Acetate, Ethyl Acetate, Petroleum, Propylene Glycol, Sodium Lauryl Sulfate, Sodium Tripolyphosphate, Terebinthina, Toluene, Xylene) 9X, 12X, 15X, 30X.

**Inactive Ingredients:**  
DeminerIALIZED water, 20% Ethanol.

**DIRECTIONS:** Adults and children 5 to 10 drops orally, 3 times 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.

LOT: XXXXXX MFD: MM/YY

## HOUSEHOLD ANTIGENS

acetone, sodium tripolyphosphate, propylene glycol, benzinum, ammonium muriaticum, sodium lauryl sulfate, terebinthina, butyl acetate, ethyl acetate, toluene, xylene, bisphenol a, petroleum, anacardium orientale, graphites, nitricum acidum, pulsatilla (pratensis), sulphur liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:44911-0536
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETONE</b> (UNII: 1364PS73AF) (ACETONE - UNII:1364PS73AF)	ACETONE	9 [hp_X] in 1 mL
<b>SODIUM TRIPOLYPHOSPHATE</b> (UNII: 5HK03SA80J) (TRIPOLYPHOSPHATE ION - UNII:5798IYA5AY)	SODIUM TRIPOLYPHOSPHATE	9 [hp_X] in 1 mL
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	9 [hp_X] in 1 mL
<b>BENZENE</b> (UNII: J64922108F) (BENZENE - UNII:J64922108F)	BENZENE	9 [hp_X] in 1 mL
<b>AMMONIUM CHLORIDE</b> (UNII: 01Q9PC255D) (AMMONIUM CATION - UNII:54S68520I4)	AMMONIUM CATION	9 [hp_X] in 1 mL
<b>SODIUM LAURYL SULFATE</b> (UNII: 368GB5141J) (LAURYL SULFATE - UNII:DIQ16UC154)	SODIUM LAURYL SULFATE	9 [hp_X] in 1 mL
<b>TURPENTINE OIL</b> (UNII: C5H0QJ6V7F) (TURPENTINE OIL - UNII:C5H0QJ6V7F)	TURPENTINE OIL	9 [hp_X] in 1 mL
<b>BUTYL ACETATE</b> (UNII: 464P5N1905) (BUTYL ACETATE - UNII:464P5N1905)	BUTYL ACETATE	9 [hp_X] in 1 mL
<b>ETHYL ACETATE</b> (UNII: 76845O8NMZ) (ETHYL ACETATE - UNII:76845O8NMZ)	ETHYL ACETATE	9 [hp_X] in 1 mL
<b>TOLUENE</b> (UNII: 3FPU23BG52) (TOLUENE - UNII:3FPU23BG52)	TOLUENE	9 [hp_X] in 1 mL
<b>XYLENE (MIXED ISOMERS)</b> (UNII: D856J1047R) (XYLENE (MIXED ISOMERS) - UNII:D856J1047R)	XYLENE (MIXED ISOMERS)	9 [hp_X] in 1 mL

<b>BISPHENOL A</b> (UNII: MLT3645I99) (BISPHENOL A - UNII:MLT3645I99)	BISPHENOL A	9 [hp_X] in 1 mL
<b>KEROSENE</b> (UNII: 1C89KKC04E) (KEROSENE - UNII:1C89KKC04E)	KEROSENE	9 [hp_X] in 1 mL
<b>SEMECARPUS ANACARDIUM JUICE</b> (UNII: Y0F0BU8RDU) (SEMECARPUS ANACARDIUM JUICE - UNII:Y0F0BU8RDU)	SEMECARPUS ANACARDIUM JUICE	12 [hp_X] in 1 mL
<b>GRAPHITE</b> (UNII: 4QQN74LH4O) (GRAPHITE - UNII:4QQN74LH4O)	GRAPHITE	12 [hp_X] in 1 mL
<b>NITRIC ACID</b> (UNII: 411VRN1TV4) (NITRIC ACID - UNII:411VRN1TV4)	NITRIC ACID	12 [hp_X] in 1 mL
<b>ANEMONE PRATENSIS</b> (UNII: 8E272251DI) (ANEMONE PRATENSIS - UNII:8E272251DI)	ANEMONE PRATENSIS	12 [hp_X] in 1 mL
<b>SULFUR</b> (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	12 [hp_X] in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44911-0536-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	03/10/2020	05/17/2028

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/10/2020	05/17/2028

**Labeler** - Energique, Inc. (789886132)

**Registrant** - Apotheca Company (844330915)

### Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0536) , api manufacture(44911-0536) , label(44911-0536) , pack(44911-0536)

Revised: 9/2024

Energique, Inc.