

**ALLERSTAT II- baptisia tinctoria, echinacea (angustifolia), allium cepa, ambrosia artemisiaefolia, arsenicum album, euphrasia officinalis, natrum sulphuricum, nux vomica, phosphorus, pulsatilla (pratensis), solidago virgaurea, 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.

Drug Facts

ACTIVE INGREDIENTS:

(in each drop): 9.98% of Allium Cepa 12X, Ambrosia Artemisiaefolia 12X, Arsenicum Album 12X, Euphrasia Officinalis 12X, Natrum Sulphuricum 12X, Phosphorus 12X, Pulsatilla (Pratensis) 12X, Solidago Virgaurea 12X, Sulphur 12X;0.10% of Baptisia Tinctoria 3X, Echinacea (Angustifolia) 3X.

INDICATIONS:

May temporarily relieve sinus congestion due to hay fever and allergic rhinitis.**

**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.

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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

ALLERSTAT II

2 fl. oz. (60 ml)

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HOMEOPATHIC REMEDY

ALLERSTAT II™

2 fl. oz. (60 ml) 20% Ethanol

Active Ingredients (in each drop):
9.98% of Allium Cepa 12X, Ambrosia Artemisiaefolia 12X, Arsenicum Album 12X, Euphrasia Officinalis 12X, Natrum Sulphuricum 12X, Nux Vomica 12X, Phosphorus 12X, Pulsatilla 12X, Solidago Virgaurea 12X, Sulphur 12X; 0.10% of Baptisia Tinctoria 3X, Echinacea 3X.

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LOT: XXXXXX
MFD: MM/YY



ALLERSTAT II

baptisia tinctoria, echinacea (angustifolia), allium cepa, ambrosia artemisiaefolia, arsenicum album,

euphrasia officinalis, natrum sulphuricum, nux vomica, phosphorus, pulsatilla (pratensis), solidago virgaurea, sulphur liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0521
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BAPTISIA TINCTORIA ROOT (UNII: 5EF0HW5WU) (BAPTISIA TINCTORIA ROOT - UNII:5EF0HW5WU)	BAPTISIA TINCTORIA ROOT	3 [hp_X] in 1 mL
ECHINACEA ANGUSTIFOLIA (UNII: VB06AV5US8) (ECHINACEA ANGUSTIFOLIA - UNII:VB06AV5US8)	ECHINACEA ANGUSTIFOLIA	3 [hp_X] in 1 mL
ONION (UNII: 492225Q21H) (ONION - UNII:492225Q21H)	ONION	12 [hp_X] in 1 mL
AMBROSIA ARTEMISIIFOLIA (UNII: 9W34L2CQ9A) (AMBROSIA ARTEMISIIFOLIA - UNII:9W34L2CQ9A)	AMBROSIA ARTEMISIIFOLIA	12 [hp_X] in 1 mL
ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) - UNII:C96613F5AV)	ARSENIC TRIOXIDE	12 [hp_X] in 1 mL
EUPHRASIA STRICTA (UNII: C9642I91WL) (EUPHRASIA STRICTA - UNII:C9642I91WL)	EUPHRASIA STRICTA	12 [hp_X] in 1 mL
SODIUM SULFATE (UNII: 0YPR65R21J) (SODIUM SULFATE ANHYDROUS - UNII:36KCS0R750)	SODIUM SULFATE	12 [hp_X] in 1 mL
STRYCHNOS NUX-VOMICA SEED (UNII: 269XH13919) (STRYCHNOS NUX-VOMICA SEED - UNII:269XH13919)	STRYCHNOS NUX-VOMICA SEED	12 [hp_X] in 1 mL
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	12 [hp_X] in 1 mL
ANEMONE PRATENSIS (UNII: 8E272251DI) (ANEMONE PRATENSIS - UNII:8E272251DI)	ANEMONE PRATENSIS	12 [hp_X] in 1 mL
SOLIDAGO VIRGAUREA FLOWERING TOP (UNII: 5405K23S50) (SOLIDAGO VIRGAUREA FLOWERING TOP - UNII:5405K23S50)	SOLIDAGO VIRGAUREA FLOWERING TOP	12 [hp_X] in 1 mL
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	12 [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:44911-0521-1	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	02/20/2020	02/11/2025

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
unapproved homeopathic		02/20/2020	02/11/2025

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0521) , api manufacture(44911-0521) , label(44911-0521) , pack(44911-0521)

Revised: 3/2022

Energique, Inc.