

JOINTOX BB- baptisia tinctoria, echinacea (angustifolia), hydrastis canadensis, borrelia burgdorferi nosode, babesia microti, ehrlichia nosode (canine), hepatitis b nosode, meningococcus nosode, arsenicum album, bryonia (alba), chelidonium majus, colchicum autumnale, kalmia latifolia, ledum palustre, lycopodium clavatum, phosphorus, rhus tox liquid Energique, Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click [here](#).

DRUG FACTS:

ACTIVE INGREDIENTS:

(in each drop): 4.75% of Hydrastis Canadensis 6X; Borrelia Burgdorferi Nosode 12X, 30X, 200X; Babesia Microti 20X, 200X, Ehrlichia Nosode (Canine) 20X, 200X; Hepatitis B Nosode 30X, 200X, Meningococcus Nosode 30X, 200X; Arsenicum Album 30C, Bryonia (Alba) 30C, Chelidonium Majus 30C, Colchicum Autumnale 30C, Kalmia Latifolia 30C, Ledum Palustre 30C, Lycopodium Clavatum 30C, Phosphorus 30C, Rhus Tox 30C. 0.10% of Baptisia Tinctoria 3X, Echinacea (Angustifolia) 3X.

INDICATIONS:

May temporarily relieve rash, fatigue, insomnia, weakness in limbs, and joint and muscle discomfort**

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

WARNINGS:

If pregnant or breastfeeding, 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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**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

JOINTOX BB

2 fl. oz. (60 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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LOT: XXXXXX MFD: MM/YY



HOMEOPATHIC REMEDY

JOINTOX BB™

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

Active Ingredients (in each drop):

4.75% of Hydrastis 6X; Borrelia Burgdorferi Nosode 12X, 30X, 200X; Babesia Microti, Ehrlichia Nosode 20X, 200X; Hepatitis B Nosode, Meningococcus Nosode 30X, 200X; Arsenicum Alb, Bryonia, Chelidonium Maj, Colchicum, Kalmia, Ledum, Lycopodium, Phos, Rhus Tox 30C. 0.10% of Baptisia, Echinacea 3X.

Inactive Ingredients:

Demineralized 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.

JOINTOX BB

baptisia tinctoria, echinacea (angustifolia), hydrastis canadensis, borrelia burgdorferi nosode, babesia microti, ehrlichia nosode (canine), hepatitis b nosode, meningococcus nosode, arsenicum album, bryonia (alba), chelidonium majus, colchicum autumnale, kalmia latifolia, ledum palustre, lycopodium clavatum, phosphorus, rhus tox liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0642
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 WHOLE (UNII: VB06AV5US8) (ECHINACEA ANGUSTIFOLIA - UNII:VB06AV5US8)		ECHINACEA ANGUSTIFOLIA WHOLE	3 [hp_X] in 1 mL
GOLDENSEAL (UNII: ZW3Z11D0JV) (GOLDENSEAL - UNII:ZW3Z11D0JV)		GOLDENSEAL	6 [hp_X] in 1 mL
BORRELIA BURGDORFERI (UNII: 0J8NV9V5Q8) (BORRELIA BURGDORFERI - UNII:0J8NV9V5Q8)		BORRELIA BURGDORFERI	12 [hp_X] in 1 mL
BABESIA MICROTI (UNII: 1948X6KEG3) (BABESIA MICROTI - UNII:1948X6KEG3)		BABESIA MICROTI	20 [hp_X] in 1 mL
EHRlichia CANIS (UNII: 970Y8T1JZY) (EHRlichia CANIS - UNII:970Y8T1JZY)		EHRlichia CANIS	20 [hp_X] in 1 mL
NEISSERIA MENINGITIDIS (UNII: V3TP2MD7F3) (NEISSERIA MENINGITIDIS - UNII:V3TP2MD7F3)		NEISSERIA MENINGITIDIS	30 [hp_X] in 1 mL
HEPATITIS B VIRUS (UNII: 77H9EM77P7) (HEPATITIS B VIRUS - UNII:77H9EM77P7)		HEPATITIS B VIRUS	30 [hp_X] in 1 mL
ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) - UNII:C96613F5AV)		ARSENIC TRIOXIDE	30 [hp_C] in 1 mL
BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)		BRYONIA ALBA ROOT	30 [hp_C] in 1 mL
CHELIDONIUM MAJUS WHOLE (UNII: 7E889U5RNN) (CHELIDONIUM MAJUS - UNII:7E889U5RNN)		CHELIDONIUM MAJUS WHOLE	30 [hp_C] in 1 mL
COLCHICUM AUTUMNALE BULB (UNII: 993QHL78E6) (COLCHICUM AUTUMNALE BULB - UNII:993QHL78E6)		COLCHICUM AUTUMNALE BULB	30 [hp_C] in 1 mL
KALMIA LATIFOLIA LEAF (UNII: 79N6542N18) (KALMIA LATIFOLIA LEAF - UNII:79N6542N18)		KALMIA LATIFOLIA LEAF	30 [hp_C] in 1 mL
LEDUM PALUSTRE TWIG (UNII: 877L01IZ0P) (LEDUM PALUSTRE TWG - UNII:877L01IZ0P)		LEDUM PALUSTRE TWG	30 [hp_C] in 1 mL
LYCOPodium CLAVATUM SPORE (UNII: C88X29Y479) (LYCOPodium CLAVATUM SPORE - UNII:C88X29Y479)		LYCOPodium CLAVATUM SPORE	30 [hp_C] in 1 mL
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)		PHOSPHORUS	30 [hp_C] in 1 mL
TOXICODENDRON PUBESCENS LEAF (UNII: 6IO182RP7A) (TOXICODENDRON PUBESCENS LEAF - UNII:6IO182RP7A)		TOXICODENDRON PUBESCENS LEAF	30 [hp_C] 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:44911-0642-1	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/26/2021	12/19/2028

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/26/2021	12/19/2028

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

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0642) , api manufacture(44911-0642) , label(44911-0642) , pack(44911-0642)

Revised: 5/2025

Energique, Inc.