

IMMUNOPAR- astragalus membranaceus, baptisia tinctoria, echinacea (angustifolia), hydrastis canadensis, lymph node (suis), medulla ossis suis, mucosa nasalis suis, spleen (suis), thymus (suis), anas barbariae, hepatis et cordis extractum, arsenicum album, lachesis mutus, phosphorus, pyrogenium 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): 9.06% of Hydrastis Canadensis 6X, Lymph Node (SUIS) 6X, Medulla Ossis Suis 6X, Mucosa Nasalis Suis 6X, Spleen (Suis) 6X, Thymus (Suis) 6X, Anas Barbariae, Hepatis ET Cordis Extractum 12X, Arsenicum Album 12X, Lachesis Mutus 12X, Phosphorus 12X, Pyrogenium 30X, 0.10% of Astragalus Membranaceus 12X, Baptisia Tinctoria 3X, Echinacea (Angustifolia) 3X

INDICATIONS:

May temporarily relieve Painful, irritated throat, mild ear disorders, and minor bronchial irritation with yellow excretion.**

**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.

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 2 days, consult your health care professional. Consult a physician for use in children under 12 years of age.

INACTIVE INGREDIENTS:

Demineralized water, 20% Ethanol.

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.

INDICATIONS:

May temporarily relieve Painful, irritated throat, mild ear disorders, and minor bronchial irritation with yellow expectoration.**

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

QUESTIONS:

Dist. by Energique, Inc.

201 Apple Blvd

Woodbine, IA 51579

800-869-8078

PACKAGE LABEL DISPLAY:

ENERGIQUE

SINCE 1987

HOMEOPATHIC REMEDY

IMMUNOPAR

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.

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

Warning - Severe or persistent sore throat or sore throat accompanied by high fever, headache, nausea, and vomiting may be serious. Consult physician promptly. Do not use more than 2 days or administer to children under 3 years of age unless directed by physician.

LOT: XXXXXX MFD: MM/YY



HOMEOPATHIC REMEDY

IMMUNOPAR™

INDICATIONS: May temporarily relieve painful, irritated throat, mild ear disorders, and minor bronchial irritation with yellow expectoration.**

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

Active Ingredients (in each drop):

9.06% of Hydrastis, Lymph Node, Medulla Ossis Suis, Mucosa Nas, Spleen, Thymus 6X; Anas Barbariae, Arsenicum Alb, Lachesis, Phos 12X; Pyrogenium 30X. 0.10% of Astragalus Membranaceus, 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 2 days, consult your health care professional. Consult a physician for use in children under 12 years of age.

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IMMUNOPAR

astragalus membranaceus, baptisia tinctoria, echinacea (angustifolia), hydrastis canadensis, lymph node (suis), medulla ossis suis, mucosa nasalis suis, spleen (suis), thymus (suis), anas barbariae, hepatis et cordis extractum, arsenicum album, lachesis mutus, phosphorus, pyrogenium liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0636
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ASTRAGALUS PROPINQUUS ROOT (UNII: 922OP8YUPF) (ASTRAGALUS PROPINQUUS ROOT - UNII:922OP8YUPF)	ASTRAGALUS PROPINQUUS ROOT	3 [hp_X] in 1 mL
BAPTISIA TINCTORIA ROOT (UNII: 5EF0HM5WU) (BAPTISIA TINCTORIA ROOT - UNII:5EF0HM5WU)	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
SUS SCROFA LYMPH (UNII: 33A7VYU29L) (SUS SCROFA LYMPH - UNII:33A7VYU29L)	SUS SCROFA LYMPH	6 [hp_X] in 1 mL
SUS SCROFA BONE MARROW (UNII: VP2CN2G7Y8) (SUS SCROFA BONE MARROW - UNII:VP2CN2G7Y8)	SUS SCROFA BONE MARROW	6 [hp_X] in 1 mL
SUS SCROFA NASAL MUCOSA (UNII: ID3Z1X61WY) (SUS SCROFA NASAL MUCOSA - UNII:ID3Z1X61WY)	SUS SCROFA NASAL MUCOSA	6 [hp_X] in 1 mL
SUS SCROFA SPLEEN (UNII: 92AMN5J79Y) (SUS SCROFA SPLEEN - UNII:92AMN5J79Y)	SUS SCROFA SPLEEN	6 [hp_X] in 1 mL
SUS SCROFA THYMUS (UNII: 7B69B0BD62) (SUS SCROFA THYMUS - UNII:7B69B0BD62)	SUS SCROFA THYMUS	6 [hp_X] in 1 mL
CAIRINA MOSCHATA HEART/LIVER AUTOLYSATE (UNII: RN2HC612GY) (CAIRINA MOSCHATA HEART/LIVER AUTOLYSATE - UNII:RN2HC612GY)	CAIRINA MOSCHATA HEART/LIVER AUTOLYSATE	12 [hp_X] in 1 mL
ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) - UNII:C96613F5AV)	ARSENIC TRIOXIDE	12 [hp_X] in 1 mL
LACHESIS MUTA VENOM (UNII: VSW71SS07I) (LACHESIS MUTA VENOM - UNII:VSW71SS07I)	LACHESIS MUTA VENOM	12 [hp_X] in 1 mL
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	12 [hp_X] in 1 mL
RANCID BEEF (UNII: 29SUH5R3HU) (RANCID BEEF - UNII:29SUH5R3HU)	RANCID BEEF	30 [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:44911-0636-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/19/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/19/2021	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0636) , api manufacture(44911-0636) , label(44911-0636) , pack(44911-0636)

Revised: 5/2021

Energique, Inc.