

NERVE PRO HP- plantago major, xanthoxylum fraxineum, colocynthis, magnesia phosphorica, agaricus muscarius, hypericum perforatum, argentum nitricum, carboneum sulphuratum, curare, thallium metallicum 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): 7.14% of Agaricus Muscarius 12X, 30X, 200X, Argentum Nitricum 30X, Carboneum Sulphuratum 30X, Colocynthis 12X, Curare 30X, Hypericum Perforatum 12X, 30X, 200X, Magnesia Phosphorica 12X, Plantago Major 6X, Thallium Metallicum 30X, Xanthoxylum Fraxineum 6X.

INDICATIONS:

May temporarily relieve symptoms of tingling, burning, and numbness in the hands and feet.**

**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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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

NERVE PRO

HP

1 fl. oz. (30 ml)

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LOT: XXXXXX MFD: MM/YY

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HOMEOPATHIC REMEDY

NERVE PRO
HP™

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

Active Ingredients (in each drop):
7.14% of Agaricus Musc 12X, 30X, 200X, Arg Nit 30X, Carboneum Sulph 30X, Colocynthis 12X, Curare 30X, Hypericum 12X, 30X, 200X, Mag Phos 12X, Plantago 6X, Thallium 30X, Xanthoxylum 6X.

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.



NERVE PRO HP

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0495
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PLANTAGO MAJOR (UNII: W2469WNO6U) (PLANTAGO MAJOR - UNII:W2469WNO6U)	PLANTAGO MAJOR	6 [hp_X] in 1 mL
ZANTHOXYLUM AMERICANUM BARK (UNII: A4KL1HMZ7T) (ZANTHOXYLUM AMERICANUM BARK - UNII:A4KL1HMZ7T)	ZANTHOXYLUM AMERICANUM BARK	6 [hp_X] in 1 mL
CITRULLUS COLOCYNTHIS FRUIT PULP (UNII: 23H32AOH17) (CITRULLUS COLOCYNTHIS FRUIT PULP - UNII:23H32AOH17)	CITRULLUS COLOCYNTHIS FRUIT PULP	12 [hp_X] in 1 mL

MAGNESIUM PHOSPHATE, DIBASIC TRIHYDRATE (UNII: HF539G9L3Q) (MAGNESIUM CATION - UNII:T6V3LHY838)	MAGNESIUM PHOSPHATE, DIBASIC TRIHYDRATE	12 [hp_X] in 1 mL
AMANITA MUSCARIA FRUITING BODY (UNII: DIF093I037) (AMANITA MUSCARIA FRUITING BODY - UNII:DIF093I037)	AMANITA MUSCARIA FRUITING BODY	12 [hp_X] in 1 mL
HYPERICUM PERFORATUM (UNII: XK4IUX8MNB) (HYPERICUM PERFORATUM - UNII:XK4IUX8MNB)	HYPERICUM PERFORATUM	12 [hp_X] in 1 mL
SILVER NITRATE (UNII: 95IT3W8JZE) (SILVER CATION - UNII:57N7B0K90A)	SILVER NITRATE	30 [hp_X] in 1 mL
CARBON DISULFIDE (UNII: S54S8B99E8) (CARBON DISULFIDE - UNII:S54S8B99E8)	CARBON DISULFIDE	30 [hp_X] in 1 mL
TUBOCURARINE CHLORIDE (UNII: 900961Z8VR) (TUBOCURARINE - UNII:W9YXS298BM)	TUBOCURARINE CHLORIDE	30 [hp_X] in 1 mL
THALLIUM (UNII: AD84R52XLF) (THALLIUM - UNII:AD84R52XLF)	THALLIUM	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:449 11-0495-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/14/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		05/14/2019	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(449 11-0495) , api manufacture(449 11-0495) , label(449 11-0495) , pack(449 11-0495)

Revised: 10/2020

Energique, Inc.