

DIZZINESS HP- calcarea carbonica, chenopodium anthelminticum, cocculus indicus, conium maculatum, nux vomica, phosphorus, pulsatilla (pratensis), silicea, tabacum, 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): 11.11% of Calcarea Carbonica 30X, Chenopodium Anthelminticum 30X, Cocculus Indicus 30X, Conium Maculatum 30X, Nux Vomica 30X, Phosphorus 30X, Pulsatilla (Pratensis) 30X, Silicea 30X, Tabacum 30X.

INDICATIONS:

May temporarily relieve symptoms of dizziness and light-headedness.**

**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 time 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 of dizziness and light-headedness.**

**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

DIZZINESS HP

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.

INDICATIONS: May temporarily relieve symptoms of dizziness and light-headedness.**

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

LOT: XXXXXX MFD: MM/YY

Dist. by Energique, Inc.
201 Apple Blvd.
Woodbine, IA 51579 **800.869.8078**



HOMEOPATHIC REMEDY

DIZZINESS HP™

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

Active Ingredients (in each drop):
11.11% of Calcarea Carbonica 30X,
Chenopodium Anthelminticum 30X,
Cocculus Indicus 30X, Conium
Maculatum 30X, Nux Vomica 30X,
Phosphorus 30X, Pulsatilla 30X,
Silicea 30X, Tabacum 30X.

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.



DIZZINESS HP

calcarea carbonica, chenopodium anthelminticum, cocculus indicus, conium maculatum, nux vomica, phosphorus, pulsatilla (pratensis), silicea, tabacum, liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:44911-0187

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OYSTER SHELL CALCIUM CARBONATE, CRUDE (UNII: 2E32821G6I) (OYSTER SHELL CALCIUM CARBONATE, CRUDE - UNII:2E32821G6I)	OYSTER SHELL CALCIUM CARBONATE, CRUDE	30 [hp_X] in 1 mL
DYSPHANIA AMBROSIOIDES (UNII: 4H5RSU087I) (CHENOPODIUM AMBROSIOIDES - UNII:4H5RSU087I)	DYSPHANIA AMBROSIOIDES	30 [hp_X] in 1 mL
ANAMIRTA COCCULUS SEED (UNII: 810258W28U) (ANAMIRTA COCCULUS SEED - UNII:810258W28U)	ANAMIRTA COCCULUS SEED	30 [hp_X] in 1 mL
CONIUM MACULATUM FLOWERING TOP (UNII: Q28R5GF371) (CONIUM MACULATUM FLOWERING TOP - UNII:Q28R5GF371)	CONIUM MACULATUM FLOWERING TOP	30 [hp_X] in 1 mL
STRYCHNOS NUX-VOMICA SEED (UNII: 269XH13919) (STRYCHNOS NUX-VOMICA SEED - UNII:269XH13919)	STRYCHNOS NUX-VOMICA SEED	30 [hp_X] in 1 mL
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	30 [hp_X] in 1 mL
PULSATILLA PRATENSIS WHOLE (UNII: 8E272251DI) (PULSATILLA PRATENSIS WHOLE - UNII:8E272251DI)	PULSATILLA PRATENSIS WHOLE	30 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	30 [hp_X] in 1 mL
TOBACCO LEAF (UNII: 6YR2608RSU) (TOBACCO LEAF - UNII:6YR2608RSU)	TOBACCO LEAF	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:44911-0187-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/08/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		07/08/2015	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
------	---------	--------	---------------------

Apotheca Company	844330915	manufacture(44911-0187) , api manufacture(44911-0187) , label(44911-0187) , pack(44911-0187)
------------------	-----------	--

Revised: 3/2022

Energique, Inc.