PNEUMO HP- ammonium muriaticum, antimonium tartaricum, arsenicum album, bryonia (alba), calcarea carbonica, lobelia inflata, natrum sulphuricum, phosphorus, spongia tosta 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 Ammonium Muriaticum 30X, Antimonium Tartaricum 30X, Arsenicum Album 30X, Bryonia (Alba) 30X, Calcarea Carbonica 30X, Lobelia Inflata 30X, Natrum Sulphuricum 30X, Phosphorus 30X, Spongia Tosta 30X.

INDICATIONS:

May temporarily relieve symptoms due to chronic lung disorders and lung inflammation.**

**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, consult your health care professional.

Consult a physician for use in children under 12 years of age.

INDICATIONS:

May temporarily relieve symptoms due to chronic lung disorders and lung inflammation.**

**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 PNEUMO HP 1 fl. oz. (30 ml)

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HOMEOPATHIC REMEDY

PNEUMO HPTM

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

Active Ingredients (in each drop): 11.11% of Ammon Mur 30X, Antimon Tart 30X, Arsenicum Alb 30X, Bryonia 30X, Calc Carb 30X, Lobelia Inf 30X, Nat Sulphuricum 30X, Phos 30X, Spongia 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.

LOT: XXXXXXX

PNEUMO HP

ammonium muriaticum, antimonium tartaricum, arsenicum album, bryonia (alba), calcarea carbonica, lobelia inflata, natrum sulphuricum, phosphorus, spongia tosta liquid

Product Information

HUMAN OTC DRUG

ORAL

Item Code (Source)

Route of Administration

Active Ingredient/Active Moiety Basis of Strength Ingredient Name Strength AMMONIUM CHLORIDE (UNII: 01Q9PC255D) (AMMONIUM CATION -30 [hp_X] AMMONIUM CATION UNII:54S6852014) in 1 mL ANTIMONY POTASSIUM TARTRATE (UNII: DL6OZ476V3) (ANTIMONY CATION ANTIMONY POTASSIUM 30 [hp_X] (3+) - UNII:069647RPT5) TARTRATE in 1 mL ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) -30 [hp X] ARSENIC TRIOXIDE UNII:C96613F5AV) in 1 mL BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT -30 [hp X] **BRYONIA ALBA ROOT** UNII:T7[046YI2B) in 1 mL OYSTER SHELL CALCIUM CARBONATE, CRUDE (UNII: 2E32821G6I) (OYSTER OYSTER SHELL CALCIUM 30 [hp_X] SHELL CALCIUM CARBONATE, CRUDE - UNII:2E32821G6I) CARBONATE, CRUDE in 1 mL 30 [hp X] LOBELIA INFLATA (UNII: 9PP1T3TC5U) (LOBELIA INFLATA - UNII: 9PP1T3TC5U) LOBELIA INFLATA in 1 mL SODIUM SULFATE (UNII: 0YPR65R21J) (SODIUM SULFATE ANHYDROUS -30 [hp X] SODIUM SULFATE UNII:36KCS0R750) in 1 mL 30 [hp_X] PHOSPHORUS PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W) in 1 mL SPONGIA OFFICINALIS SKELETON, ROASTED (UNII: 1PIP394IID) (SPONGIA SPONGIA OFFICINALIS 30 [hp_X] OFFICINALIS SKELETON, ROASTED - UNII:1PIP394IID) SKELETON, ROASTED in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

# It	em Code	Package Description	Marketing Start Date	Marketing End Date
		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/13/2016	03/30/2025

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
unapproved homeopathic		06/17/2015	03/30/2025

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment				
Name	Address	ID/FEI	Business Operations	
Apotheca Company		844330915	manufacture(44911-0169) , api manufacture(44911-0169) , label(44911-0169) , pack(44911-0169)	

Revised: 3/2022

Energique, Inc.