BREATH 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 productive cough, chest congestion, shortness of breath and Fatigue.**

**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 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 productive cough, chest congestion, shortness of breath and Fatigue.**

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

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HOMEOPATHIC REMEDY

BREATH HP[™]

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: XXXXXX

BREATH HP

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

Product Information

Product	Туре
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HUMAN OTC DRUG

Item Code (Source)

Route of Administration ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
AMMONIUM CHLORIDE (UNII: 01Q9PC255D) (AMMONIUM CATION - UNII:54S68520I4)	AMMONIUM CATION	30 [hp_X] in 1 mL	
ANTIMONY POTASSIUM TARTRATE (UNII: DL6OZ476V3) (ANTIMONY CATION (3+) - UNII:069647RPT5)	ANTIMONY POTASSIUM TARTRATE	30 [hp_X] in 1 mL	
ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) - UNII:C96613F5AV)	ARSENIC TRIOXIDE	30 [hp_X] in 1 mL	
BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)	BRYONIA ALBA ROOT	30 [hp_X] in 1 mL	
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	
LOBELIA INFLATA (UNII: 9PP1T3TC5U) (LOBELIA INFLATA - UNII:9PP1T3TC5U)	LOBELIA INFLATA	30 [hp_X] in 1 mL	
SODIUM SULFATE (UNII: 0YPR65R21J) (SODIUM SULFATE ANHYDROUS - UNII: 36KCS0R750)	SODIUM SULFATE	30 [hp_X] in 1 mL	
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W)	PHOSPHORUS	30 [hp_X] in 1 mL	
SPONGIA OFFICINALIS SKELETON, ROASTED (UNII: 1PIP394IID) (SPONGIA OFFICINALIS SKELETON, ROASTED - UNII:1PIP394IID)	SPONGIA OFFICINALIS SKELETON, ROASTED	30 [hp_X] in 1 mL	

Inactive Ingredients

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

Packaging

	#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:44911- 0591-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	06/01/2021	
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Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
unapproved homeopathic		06/01/2021	

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0591) , api manufacture(44911-0591) , label(44911-0591) , pack(44911-0591)

Revised: 6/2021

Energique, Inc.