

**FUNGUSTAT II- echinacea (angustifolia), fucus vesiculosus, hydrastis canadensis, phytolacca decandra, kreosotum, natrum muriaticum, candida albicans, lycopodium clavatum, pulsatilla (pratensis), sepia 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): 11.09% of Candida Albicans 12X, 30X, Hydrastis Canadensis 6X, Kreosotum 12X, Lycopodium Clavatum 30X, Natrum Muriaticum 12X, Phytolacca Decandra 6X, Pulsatilla (Pratensis) 30X, Sepia 30X; 0.10% of Echinacea (Angustifolia) 3X, Fucus Vesiculosus 3X.

INDICATIONS:

May temporarily relieve bloating of the stomach, flatulence and rhinitis.**

**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.

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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

FUNGUSTAT II

1 fl. oz. (30 ml)

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LOT: XXXXXX MFD: MM/YY

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HOMEOPATHIC REMEDY

FUNGUSTAT II™

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

Active Ingredients (in each drop):
11.09% of Candida Alb 12X, 30X, Hydrastis 6X, Kreosotum 12X, Lycopodium 30X, Nat Mur 12X, Phytolacca 6X, Pulsatilla 30X, Sepia 30X; 0.10% of Echinacea 3X, Fucus 3X.

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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.



FUNGUSTAT II

echinacea (angustifolia), fucus vesiculosus, hydrastis canadensis, phytolacca decandra, kreosotum, natrum muriaticum, candida albicans, lycopodium clavatum, pulsatilla (pratensis), sepia liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:44911-0437
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ECHINACEA ANGUSTIFOLIA (UNII: VB06AV5US8) (ECHINACEA ANGUSTIFOLIA - UNII:VB06AV5US8)	ECHINACEA ANGUSTIFOLIA	3 [hp_X] in 1 mL
FUCUS VESICULOSUS (UNII: 535G2ABX9M) (FUCUS VESICULOSUS - UNII:535G2ABX9M)	FUCUS VESICULOSUS	3 [hp_X] in 1 mL
GOLDENSEAL (UNII: ZW3Z11D0JV) (GOLDENSEAL - UNII:ZW3Z11D0JV)	GOLDENSEAL	6 [hp_X] in 1 mL
PHYTOLACCA AMERICANA ROOT (UNII: 11E6VI8VEG) (PHYTOLACCA AMERICANA ROOT - UNII:11E6VI8VEG)	PHYTOLACCA AMERICANA ROOT	6 [hp_X] in 1 mL
WOOD CREOSOTE (UNII: 3JYG22FD73) (WOOD CREOSOTE - UNII:3JYG22FD73)	WOOD CREOSOTE	12 [hp_X] in 1 mL
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	12 [hp_X] in 1 mL
CANDIDA ALBICANS (UNII: 4D7G21HDBC) (CANDIDA ALBICANS - UNII:4D7G21HDBC)	CANDIDA ALBICANS	12 [hp_X] in 1 mL
LYCOPODIUM CLAVATUM SPORE (UNII: C88X29Y479) (LYCOPODIUM CLAVATUM SPORE - UNII:C88X29Y479)	LYCOPODIUM CLAVATUM SPORE	30 [hp_X] in 1 mL
ANEMONE PRATENSIS (UNII: 8E272251DI) (ANEMONE PRATENSIS - UNII:8E272251DI)	ANEMONE PRATENSIS	30 [hp_X] in 1 mL
SEPIA OFFICINALIS JUICE (UNII: QDL83WN8C2) (SEPIA OFFICINALIS JUICE - UNII:QDL83WN8C2)	SEPIA OFFICINALIS JUICE	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:44911-0437-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/05/2017	04/13/2025

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/05/2017	04/13/2025

Labeler - Energique, Inc. (789886132)

Registrant - Apotheca Company (844330915)**Establishment**

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(44911-0437) , api manufacture(44911-0437) , label(44911-0437) , pack(44911-0437)

Revised: 3/2022

Energique, Inc.