

IGNATIA BRYOPHYLLUM- ignatia bryophyllum liquid

Uriel Pharmacy 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.

Ignatia Bryophyllum Special Order

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops. Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredients: Bryophyllum (Life plant) 3X, Ignatia (St. Ignatius' bean) 4X, Lachesis (Bushmaster venom) 12X, Argentum (Silver) 10/20/30X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol

Use: Temporary relief of hot flashes.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858

Made with care by Uriel, East Troy, WI 53120

www.urielpharmacy.com

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**Ignatia
Bryophyllum
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Homeopathic Liquid
net vol. 2 fl. oz (60ml)

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www.urielpharmacy.com Lot:

IGNATIA BRYOPHYLLUM

ignatia bryophyllum liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8306
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
STRYCHNOS IGNATII SEED (UNII: 1NM3M2487K) (STRYCHNOS IGNATII SEED - UNII:1NM3M2487K)	STRYCHNOS IGNATII SEED	4 [hp_X] in 1 mL
LACHESIS MUTA VENOM (UNII: VSW71SS07I) (LACHESIS MUTA VENOM - UNII:VSW71SS07I)	LACHESIS MUTA VENOM	12 [hp_X] in 1 mL
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)	SILVER	10 [hp_X] in 1 mL
KALANCHOE DAIGREMONTIANA LEAF (UNII: L6X13JKL8O) (KALANCHOE DAIGREMONTIANA LEAF - UNII:L6X13JKL8O)	KALANCHOE DAIGREMONTIANA LEAF	3 [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:48951-8306-3	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)**Establishment**

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-8306)

Revised: 4/2018

Uriel Pharmacy Inc.