

REYNOUTRIA LYME- reynoutria lyme 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.

Reynoutria Lyme

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: Reynoutria japonica (Japanese knotweed) 3X, Dipsacus sylvestris (Teasel) 4X, Helleborus (Christmas rose) 4X, Astragalus membranaceus e rad. (Milk vetch) 6X, Argentum (Silver) 17X, Cuprum (Copper) 17X, Quartz (Rock crystal) 17X, Apis (Honeybee) 21X, Astragalus membranaceus e fol. et sem. (Milk vetch) 30X, Aurum (Metallic gold) 30X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol

Use: Immune support.

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

shopuriel.com Lot:

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Lot:


**Reynoutria
Lyme**
Homeopathic Liquid
net vol. 2 fl. oz (60ml)

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REYNOUTRIA LYME

reynoutria lyme liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8396
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
COPPER (UNII: 789U1901C5) (COPPER - UNII:789U1901C5)	COPPER	17 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	17 [hp_X] in 1 mL
APIS MELLIFERA (UNII: 7S82P3R43Z) (APIS MELLIFERA - UNII:7S82P3R43Z)	APIS MELLIFERA	21 [hp_X] in 1 mL
ASTRAGALUS CHINENSIS SEED (UNII: U046B39ADE) (ASTRAGALUS CHINENSIS SEED - UNII:U046B39ADE)	ASTRAGALUS CHINENSIS SEED	30 [hp_X] in 1 mL
GOLD (UNII: 79Y1949PYO) (GOLD - UNII:79Y1949PYO)	GOLD	30 [hp_X] in 1 mL
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)	SILVER	17 [hp_X] in 1 mL
HELLEBORUS NIGER ROOT (UNII: 608DGJ6815) (HELLEBORUS NIGER ROOT - UNII:608DGJ6815)	HELLEBORUS NIGER ROOT	4 [hp_X] in 1 mL
DIPSACUS FULLONUM WHOLE (UNII: 1FT87R544S) (DIPSACUS FULLONUM WHOLE - UNII:1FT87R544S)	DIPSACUS FULLONUM WHOLE	4 [hp_X] in 1 mL
ASTRAGALUS PROPINQUUS ROOT (UNII: 922OP8YUPF) (ASTRAGALUS PROPINQUUS ROOT - UNII:922OP8YUPF)	ASTRAGALUS PROPINQUUS ROOT	6 [hp_X] in 1 mL
REYNOUTRIA JAPONICA LEAF (UNII: 2540B7G25G) (REYNOUTRIA JAPONICA LEAF - UNII:2540B7G25G)	REYNOUTRIA JAPONICA LEAF	3 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8396-3	60 mL in 1 BOTTLE, GLASS; 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(48951-8396)

Revised: 10/2022

Uriel Pharmacy, Inc.