

BERBERIS LARIX- berberis larix 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.

Berberis Larix

Directions: FOR ORAL USE.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredients: Achillea (Yarrow) 3X, Berberis (Barberry) 3X, Equisetum (Common horsetail) 3X, Cantharis (Spanish fly) 6X, Apis (Honeybee) 8X, Pyrite (Nat. Iron disulfide) 8X, Resina laricis (Larch resin) 8X, Vesica urinaria (Bovine urinary bladder) 8X

Inactive Ingredients: Water, Salt, Lactose

Use: Temporary relief of urinary irritation.

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. Contains traces of lactose. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use.

Questions? Call 866.642.2858 Made by Uriel, East Troy, WI 53120 www.urielpharmacy.com

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www.urielpharmacy.com Lot:



Berberis Larix

Homeopathic Ampules
 net vol. 0.3 fl. oz (10 x 1 ml)

Berberis Larix

BERBERIS LARIX			
berberis larix liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-2042
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ACHILLEA MILLEFOLIUM (UNII: 2FXJ6SW4PK) (ACHILLEA MILLEFOLIUM -		ACHILLEA MILLEFOLIUM	3 [hp_X]

UNII:2FXJ6SW4PK)	ACHILLEA MELLEFOLIUM	in 1 mL
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	3 [hp_X] in 1 mL
EQUISETUM ARVENSE TOP (UNII: 1DP6Y6B65Z) (EQUISETUM ARVENSE TOP - UNII:1DP6Y6B65Z)	EQUISETUM ARVENSE TOP	3 [hp_X] in 1 mL
LYTTA VESICATORIA (UNII: 3Q034RO3BT) (LYTTA VESICATORIA - UNII:3Q034RO3BT)	LYTTA VESICATORIA	6 [hp_X] in 1 mL
APIS MELLIFERA (UNII: 7S82P3R43Z) (APIS MELLIFERA - UNII:7S82P3R43Z)	APIS MELLIFERA	8 [hp_X] in 1 mL
FERROUS DISULFIDE (UNII: 132N09W4PR) (FERROUS CATION - UNII:GW89581OWR)	FERROUS CATION	8 [hp_X] in 1 mL
LARIX DECIDUA RESIN (UNII: AD8LJ73GQF) (LARIX DECIDUA RESIN - UNII:AD8LJ73GQF)	LARIX DECIDUA RESIN	8 [hp_X] in 1 mL
SUS SCROFA URINARY BLADDER (UNII: 3G7U72W8DA) (SUS SCROFA URINARY BLADDER - UNII:3G7U72W8DA)	SUS SCROFA URINARY BLADDER	8 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LACTOSE (UNII: J2B2A4N98G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-2042-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package		

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(48951-2042)

Revised: 5/2018

Uriel Pharmacy Inc.