LACHESIS BELLADONNA- laches is belladonna 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.

Laches is Belladonna

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: Mercurialis (Dog's mercury) 3X, Atropa belladonna (Nightshade) 4X, Hepar sulfuris (Liver of lime) 6X, Lachesis e veneno (Bushmaster venom) 12X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol, Lactose

Use: Promotes healing of wounds.

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. Contains traces of lactose. 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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Homeopathic Liquid net vol. 2 fl. oz (60ml)

Belladonna

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LACHESIS BELLADONNA

lachesis belladonna liquid

Product Information

1 Todact Into mation			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-6013
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
MERCURIALIS PERENNIS (UNII: Q35465A1MA) (MERCURIALIS PERENNIS - UNII:Q35465A1MA)	MERCURIALIS PERENNIS	3 [hp_X] in 1 mL	

ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII:WQZ3G9PF0H)	ATROPA BELLADONNA	4 [hp_X] in 1 mL
CALCIUM SULFIDE (UNII: 1MBW07J51Q) (CALCIUM SULFIDE - UNII:1MBW07J51Q)	CALCIUM SULFIDE	6 [hp_X] in 1 mL
LACHESIS MUTA VENOM (UNII: VSW71SS07I) (LACHESIS MUTA VENOM - UNII: VSW71SS07I)	LACHESIS MUTA VENOM	12 [hp_X] in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
ALCOHOL (UNII: 3K9958V90M)			
LACTOSE (UNII: J2B2A4N98G)			

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:48951-6013- 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(48951-6013)	

Revised: 4/2018 Uriel Pharmacy Inc.