ACONITUM BRYONIA- aconitum bryonia 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.

Aconitum Bryonia

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: Aconitum (Monkshood) 3X, Bryonia (White bryony) 3X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol

Use: Temporary relief of flu symptoms.

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

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Homeopathic Liquid net vol. 2 fl. oz (60ml) Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if alergic to any ingredient. C doctor before use for serious conditions or if conditions worsen or persit. If pregnant or number consult a doctor before use. Do not use if safety seal braken or mixing.

Questions? Call 866,642,2058
Made with care by Uhiel, East Tray, WI 53120

ACONITUM BRYONIA

aconitum bryonia liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-1359	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACONITUM NAPELLUS (UNII: U0NQ8555JD) (ACONITUM NAPELLUS - UNII:U0NQ8555JD)	ACONITUM NAPELLUS	3 [hp_X] in 1 mL	
BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)	BRYONIA ALBA ROOT	3 [hp_X] in 1 mL	

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

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:48951- 1359-3	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	

Monograph Marketing Start Marketing En Date Date
09/01/2009

Labeler - Uriel Pharmacy, Inc. (043471163)

Establishment				
Name	Address	ID/FEI	Business Operations	
Uriel Pharmacy, Inc.		043471163	manufacture(48951-1359)	

Revised: 11/2022 Uriel Pharmacy, Inc.