

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, 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 www.urielpharmacy.com

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**Aconitum
Bryonia**
Homeopathic Liquid
net vol. 2 fl. oz (60ml)

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.
Lot:

ACONITUM BRYONIA

aconitum bryonia liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-9-170
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: T7J046 YI2B) (BRYONIA ALBA ROOT - UNII:T7J046 YI2B)	BRYONIA ALBA ROOT	3 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-9170-3	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 0: Not a Combination Product		

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(48951-9170)

Revised: 4/2018

Uriel Pharmacy Inc.