BRYONIA - bryonia liquid Newton Laboratories, 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.

Bryonia

INDICATIONS & USAGE SECTION

Liver Remedy, Abscess, Joint Pain, Aches & Pains, Headache, Sore Throat, Shortness of Breath, Cough.

DOSAGE & ADMINISTRATION SECTION

Directions: Ages 12 and up, take 6 drops by mouth (ages 0 to 11, give 3 drops) as needed or as directed by a health professional. Sensitive persons begin with 1 drop and gradually increase to full dose.

OTC - ACTIVE INGREDIENT SECTION

Bryonia 15x, 10x, 200c, 30c.

PURPOSE SECTION

Liver Remedy, Abscess, Joint Pain, Aches & Pains, Headache, Sore Throat, Shortness of Breath, Cough.

INACTIVE INGREDIENT SECTION

Inactive Ingredients: USP Purified Water; USP Gluten-free, non-GMO, organic cane alcohol 20%.

OTC - QUESTIONS SECTION

www.newtonlabs.net Newton Laboratories, Inc. FDA Est # 1051203 - Conyers, GA 30012 Questions? 1.800.448.7256

WARNINGS SECTION

Warning: Keep out of reach of children. Do not use if tamper-evident seal is broken or missing. If symptoms worsen or persist for more than a few days, consult a doctor. If **pregnant or breast-feeding**, ask a doctor before use.

OTC - PREGNANCY OR BREAST FEEDING SECTION

Consult a licensed healthcare professional if pregnant, nursing or if symptoms worsen or persist for more than a few days.

OTC - KEEP OUT OF REACH OF CHILDREN SECTION

Keep out of reach of children.

PACKAGE LABEL



BRYONIA

bryonia liquid

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55714-6107
Route of Administration	ORAL		

Active Ingredient/Active Meiety

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Ingredient Name	Basis of Strength	Strength	
Bryonia Alba Root (UNII: T7J046 YI2B) (Bryonia Alba Root - UNII:T7J046 YI2B)	Bryonia Alba Root	15 [hp_X] in 1 mL	

Inactive Ingredients

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Ingredient Name	Strength			
Alcohol (UNII: 3K9958V90M)				
Water (UNII: 059QF0KO0R)				

Packaging

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ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:55714-6107-1	30 mL in 1 BOTTLE, GLASS		

Marketing Information

The first matter			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2011	

Labeler - Newton Laboratories, Inc. (788793610)

Registrant - Newton Laboratories, Inc. (788793610)

Establishment				
Name	Address	ID/FEI	Business Operations	
Newton Laboratories, Inc.		788793610	MANUFACTURE(55714-6107)	

Revised: 9/2011 Newton Laboratories, Inc.