

**ASTRAGALUS ARNICA- astragalus arnica 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.*

**Astragalus Arnica**

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: Astragalus (Milk vetch) 3X, Arnica 30X

Inactive Ingredients: Distilled water, Organic cane alcohol

Use: Temporary relief of headache.

**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

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: Astragalus (Milk vetch) 3X, Arnica 30X  
 Inactive Ingredients: Distilled water, 20% Organic cane alcohol  
 Use: Temporary relief of headache.  
 Lot:



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<b>ASTRAGALUS ARNICA</b>			
astragalus arnica liquid			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:48951-1171
<b>Route of Administration</b>	ORAL		
<b>Active Ingredient/Active Moiety</b>			

Ingredient Name	Basis of Strength	Strength
ASTRAGALUS PROPINQUUS ROOT (UNII: 922OP8 YUPF) (ASTRAGALUS PROPINQUUS ROOT - UNII:922OP8 YUPF)	ASTRAGALUS PROPINQUUS ROOT	3 [hp_X] in 1 mL
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	30 [hp_X] in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
ALCOHOL (UNII: 3K9958V90M)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-1171-3	60 mL in 1 BOTTLE, GLASS; 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(48951-1171)

Revised: 4/2018

Uriel Pharmacy Inc.