

ASTRAGALUS FORMICA- astragalus formica 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 Formica

Directions: FOR ORAL USE.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredients: Arnica 3X, Astragalus (Milk vetch) 3X, Formica (Red wood ant) 6X, Argentum (Silver) 30X

Inactive Ingredients: Water, Salt

"prepared using rhythmical processes"

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.

Questions? Call 866.642.2858

Made by Uriel, East Troy, WI 53120

shopuriel.com Lot:

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Made by Uriel, East Troy, WI 53120
shopuriel.com Lot:

uriel

**Astragalus
Formica**

Homeopathic Ampules
net vol. 0.3 fl. oz (10 x 1 ml)

Astragalus Formica

ASTRAGALUS FORMICA

astragalus formica liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:48951-1174

Route of Administration		ORAL		
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ARNICA MONTANA (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)		ARNICA MONTANA	3 [hp_X] in 1 mL	
ASTRAGALUS PROPINQUUS ROOT (UNII: 922OP8YUPF) (ASTRAGALUS PROPINQUUS ROOT - UNII:922OP8YUPF)		ASTRAGALUS PROPINQUUS ROOT	3 [hp_X] in 1 mL	
FORMICA RUFA (UNII: 55H0W83JO5) (FORMICA RUFA - UNII:55H0W83JO5)		FORMICA RUFA	6 [hp_X] in 1 mL	
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)		SILVER	30 [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:48951-1174-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package		
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(48951-1174)