

**BAMBUSA RHUS- bambusa rhus pellet**  
**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.*

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**Bambusa Rhus**

Directions: FOR ORAL USE ONLY.

Dissolve pellets under the tongue 3-4 times daily. Ages 12 and older: 10 pellets. Ages 2-11: 5 pellets. Under age 2: Consult a doctor.

Active Ingredients: Gelsemium (Yellow jasmine) 4X, Gnaphalium (Edelweiss) 4X, Aconitum (Monkshood) 6X, Bambusa (Bamboo) 6X, Betonica (Wood betony) 6X, Mandragora (Mandrake) 6X, Rhus tox. (Sumac) 6X, Rosmarinus (Rosemary) 6X, Formica (Red wood ant) 7X, Disci intervert. (Bovine intervertebral discs of the cerv. thor. and lumbar spine) 8X, Granite (Primitive rock containing quartz, feldspar and mica) 10X, Jasper (Semiprecious stone) 12X, Argentum (Silver) 20X, Arnica 20X

Inactive Ingredient: Sucrose

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. Contains sugar. Diabetics and persons intolerant of sucrose (sugar): Consult a doctor before use. 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  
 Uriel, East Troy, WI 53120  
[www.urielpharmacy.com](http://www.urielpharmacy.com)

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<b>BAMBUSA RHUS</b>			
bambusa rhus pellet			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:48951-2017
<b>Route of Administration</b>	ORAL		
<b>Active Ingredient/Active Moiety</b>			

Ingredient Name	Basis of Strength	Strength
<b>GELSEMIUM SEMPERVIRENS ROOT</b> (UNII: 639KR60Q1Q) (GELSEMIUM SEMPERVIRENS ROOT - UNII:639KR60Q1Q)	GELSEMIUM SEMPERVIRENS ROOT	4 [hp_X]
<b>PSEUDOGNAPHALIUM LUTEOALBUM LEAF</b> (UNII: BGI20Z6M57) (PSEUDOGNAPHALIUM LUTEOALBUM LEAF - UNII:BGI20Z6M57)	PSEUDOGNAPHALIUM LUTEOALBUM LEAF	4 [hp_X]
<b>ACONITUM NAPELLUS</b> (UNII: U0NQ8555JD) (ACONITUM NAPELLUS - UNII:U0NQ8555JD)	ACONITUM NAPELLUS	6 [hp_X]
<b>BAMBUSA VULGARIS WHOLE</b> (UNII: WCD45M1BSK) (BAMBUSA VULGARIS WHOLE - UNII:WCD45M1BSK)	BAMBUSA VULGARIS WHOLE	6 [hp_X]
<b>STACHYS OFFICINALIS</b> (UNII: UO9989Y17N) (STACHYS OFFICINALIS - UNII:UO9989Y17N)	STACHYS OFFICINALIS	6 [hp_X]
<b>MANDRAGORA OFFICINARUM ROOT</b> (UNII: I2XCB174VB) (MANDRAGORA OFFICINARUM ROOT - UNII:I2XCB174VB)	MANDRAGORA OFFICINARUM ROOT	6 [hp_X]
<b>TOXICODENDRON PUBESCENS LEAF</b> (UNII: 6IO182RP7A) (TOXICODENDRON PUBESCENS LEAF - UNII:6IO182RP7A)	TOXICODENDRON PUBESCENS LEAF	6 [hp_X]
<b>ROSMARINUS OFFICINALIS FLOWERING TOP</b> (UNII: 8JM482TI79) (ROSMARINUS OFFICINALIS FLOWERING TOP - UNII:8JM482TI79)	ROSMARINUS OFFICINALIS FLOWERING TOP	6 [hp_X]
<b>FORMICA RUFA</b> (UNII: 55H0W83JO5) (FORMICA RUFA - UNII:55H0W83JO5)	FORMICA RUFA	7 [hp_X]
<b>BOS TAURUS INTERVERTEBRAL DISC</b> (UNII: 0A04Z76C13) (BOS TAURUS INTERVERTEBRAL DISC - UNII:0A04Z76C13)	BOS TAURUS INTERVERTEBRAL DISC	8 [hp_X]
<b>PENOXSULAM</b> (UNII: 784ELC1SCZ) (PENOXSULAM - UNII:784ELC1SCZ)	PENOXSULAM	10 [hp_X]
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	12 [hp_X]
<b>SILVER</b> (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)	SILVER	20 [hp_X]
<b>ARNICA MONTANA</b> (UNII: O80TY208ZW) (ARNICA MONTANA - UNII:O80TY208ZW)	ARNICA MONTANA	20 [hp_X]

### Inactive Ingredients

Ingredient Name	Strength
<b>SUCROSE</b> (UNII: C151H8M554)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	3mm
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-2017-2	1350 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	

## Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-2017)

Revised: 5/2018

Uriel Pharmacy Inc.