BRIDGED RING COMBINATION 9311- bridged ring combination liquid Professional Complementary Health Formulas

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.

R311

ACTIVE INGREDIENTS

Cimicifuga racemosa 3X
Gelsemium sempervirens 3X
Natrum carbonicum 3X
Endosulfan 12X
Heptachlor 12X
Silicea 12X
Thuja occidentalis 12X

QUESTIONS

Professional Formulas
PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of occasional headache, nausea or vomiting, confusion, or weakness.*

*Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

WARNINGS

Consult a doctor if condition worsens or if symptoms persist. Keep out of the reach of children. In case of overdose, get medical help or contact a poison control center right away. If pregnant or breastfeeding, ask a healthcare professional before use.

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DIRECTIONS

Place drops under tongue 30 minutes before/after meals. Adults and children 12 years and over: Take 10 drops up to 3 times per day. Consult a physician for use in children under 12 years of age.

OTHER INFORMATION

Tamper resistant. If seal is broken, do not use. After opening, close container tightly and store at room temperature away from heat.

INACTIVE INGREDIENTS

40% ethanol, purified water.

LABEL

Est 1985

Professional Formulas

Complementary Health

Bridged Ring Combination

Homeopathic Remedy

1 FL. OZ. (29.5 mL)



BRIDGED RING COMBINATION 9311

bridged ring combination liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-9311
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength

BLACK COHOSH (UNII: K73E24S6X9) (BLACK COHOSH - UNII:K73E24S6X9)	BLACK COHOSH	3 [hp_X] in 29.5 mL
GELSEMIUM SEMPERVIRENS ROOT (UNII: 639KR60Q1Q) (GELSEMIUM SEMPERVIRENS ROOT - UNII:639KR60Q1Q)	GELSEMIUM SEMPERVIRENS ROOT	3 [hp_X] in 29.5 mL
SODIUM CARBONATE (UNII: 45P3261C7T) (CARBONATE ION - UNII:7UJQ50PE7D)	SODIUM CARBONATE	3 [hp_X] in 29.5 mL
Malathion (UNII: U5N7SU872W) (MALATHION - UNII:U5N7SU872W)	Malathion	12 [hp_X] in 29.5 mL
HEPTACHLOR (UNII: 7GLS9ACN3L) (HEPTACHLOR - UNII:7GLS9ACN3L)	HEPTACHLOR	12 [hp_X] in 29.5 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	12 [hp_X] in 29.5 mL
THUJA OCCIDENTALIS LEAF (UNII: 0T0DQN8786) (THUJA OCCIDENTALIS LEAF - UNII:0T0DQN8786)	THUJA OCCIDENTALIS LEAF	12 [hp_X] in 29.5 mL

Inactive Ingredients	
Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
			29.5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	08/15/1985	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/15/1984	

Labeler - Professional Complementary Health Formulas (167339027)

Registrant - Natural Pharmaceutical Manufacturing LLC (015624923)

Establishment				
Name	Address	ID/FEI	Business Operations	
Natural Pharmaceutical Manufacturing LLC		015624923	manufacture(63083-9311)	

Revised: 8/2019 Professional Complementary Health Formulas