

## **CHLORINATED RING COMBINATION 9314- chlorinated 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.*

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**R314**

### **ACTIVE INGREDIENTS**

Cimicifuga racemosa 3X  
Gelsemium sempervirens 3X  
Nux vomica 6X  
Berberis vulgaris 10X  
Chelidonium majus 10X  
2,4,5 T Ester 12X  
Hexachlorbenzol 14X  
Pentachlorophenol 14X  
Polychlorinated biphenyls (PCBs) 14X

### **QUESTIONS**

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

### **INDICATIONS**

For the temporary relief of occasional headache, upper respiratory irritation, nausea or vomiting, red or irritated skin or eyes, 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  
Chlorinated Ring Combination  
Homeopathic Remedy  
1 FL. OZ. (29.5 mL)



CHLORINATED RING COMBINATION 9314			
chlorinated ring combination liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-9314
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
<b>BLACK COHOSH</b> (UNII: K73E24S6X9) (BLACK COHOSH - UNII:K73E24S6X9)	BLACK COHOSH	3 [hp_X] in 29.5 mL
<b>GELSEMIUM SEMPERVIRENS ROOT</b> (UNII: 639KR60Q1Q) (GELSEMIUM SEMPERVIRENS ROOT - UNII:639KR60Q1Q)	GELSEMIUM SEMPERVIRENS ROOT	3 [hp_X] in 29.5 mL
<b>STRYCHNOS NUX-VOMICA SEED</b> (UNII: 269XH13919) (STRYCHNOS NUX-VOMICA SEED - UNII:269XH13919)	STRYCHNOS NUX-VOMICA SEED	6 [hp_X] in 29.5 mL
<b>BERBERIS VULGARIS ROOT BARK</b> (UNII: 1TH8Q20J0U) (BERBERIS VULGARIS ROOT BARK - UNII:1TH8Q20J0U)	BERBERIS VULGARIS ROOT BARK	10 [hp_X] in 29.5 mL
<b>CHELIDONIUM MAJUS WHOLE</b> (UNII: 7E889U5RNN) (CHELIDONIUM MAJUS WHOLE - UNII:7E889U5RNN)	CHELIDONIUM MAJUS WHOLE	10 [hp_X] in 29.5 mL
<b>ACETIC ACID, (2,4,5-TRICHLOROPHENOXY)-, 2-ETHYL-4-METHYLPENTYL ESTER</b> (UNII: 03HP3QF78W) (ACETIC ACID, (2,4,5-TRICHLOROPHENOXY)-, 2-ETHYL-4-METHYLPENTYL ESTER - UNII:03HP3QF78W)	ACETIC ACID, (2,4,5-TRICHLOROPHENOXY)-, 2-ETHYL-4-METHYLPENTYL ESTER	12 [hp_X] in 29.5 mL
<b>HEXACHLORO BENZENE</b> (UNII: 4Z87H0LKUY) (HEXACHLORO BENZENE - UNII:4Z87H0LKUY)	HEXACHLORO BENZENE	14 [hp_X] in 29.5 mL
<b>PENTACHLOROPHENOL</b> (UNII: D9BSU0SE4T) (PENTACHLOROPHENOL - UNII:D9BSU0SE4T)	PENTACHLOROPHENOL	14 [hp_X] in 29.5 mL
<b>FENSON</b> (UNII: DFC2HB4I0K) (FENSON - UNII:DFC2HB4I0K)	FENSON	14 [hp_X] in 29.5 mL

Inactive Ingredients	
Ingredient Name	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>WATER</b> (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63083-9314-1	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(63083-9314)

