

## **VIROX 6022- virox 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.*

-----

**X22**

### **ACTIVE INGREDIENTS**

Cinchona officinalis 3X  
Nux moschata 3X  
Gelsemium sempervirens 6X  
Pulsatilla 6X  
Sepia 6X  
Crotalus horridus 8X  
Lachesis mutus 8X

### **QUESTIONS**

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

### **INDICATIONS**

For the temporary relief of runny nose, sneezing, cough, congestion, occasional headache, chills, sweats, sore throat, minor joint or muscle aches or pains, or exhaustion.\*

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

### **WARNINGS**

Persistent symptoms may be a sign of a serious condition. If symptoms persist or are accompanied by a fever, rash, or persistent headache, consult a doctor. 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.

Keep out of the reach of children.

If pregnant or breastfeeding, ask a healthcare professional before use.

### **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**

20% ethanol, purified water.

**LABEL**

Est 1985  
Professional Formulas  
Complementary Health  
Virox  
Homeopathic Remedy  
2 FL. OZ. (59 mL)



VIROX 6022			
virox liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-6022
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
<b>CINCHONA OFFICINALIS BARK</b> (UNII: S003A158SB) (CINCHONA OFFICINALIS BARK - UNII:S003A158SB)	CINCHONA OFFICINALIS BARK	3 [hp_X] in 59 mL
<b>NUTMEG</b> (UNII: AEE24M3MQ9) (NUTMEG - UNII:AEE24M3MQ9)	NUTMEG	3 [hp_X] in 59 mL
<b>GELSEMIUM SEMPERVIRENS ROOT</b> (UNII: 639KR60Q1Q) (GELSEMIUM SEMPERVIRENS ROOT - UNII:639KR60Q1Q)	GELSEMIUM SEMPERVIRENS ROOT	6 [hp_X] in 59 mL
<b>PULSATILLA MONTANA WHOLE</b> (UNII: 24K790T39B) (PULSATILLA MONTANA WHOLE - UNII:24K790T39B)	PULSATILLA MONTANA WHOLE	6 [hp_X] in 59 mL
<b>SEPIA OFFICINALIS JUICE</b> (UNII: QDL83WN8C2) (SEPIA OFFICINALIS JUICE - UNII:QDL83WN8C2)	SEPIA OFFICINALIS JUICE	6 [hp_X] in 59 mL
<b>CROTALUS HORRIDUS HORRIDUS VENOM</b> (UNII: YHA2XLJ956) (CROTALUS HORRIDUS HORRIDUS VENOM - UNII:YHA2XLJ956)	CROTALUS HORRIDUS HORRIDUS VENOM	8 [hp_X] in 59 mL
<b>LACHESIS MUTA VENOM</b> (UNII: VSW71SS07I) (LACHESIS MUTA VENOM - UNII:VSW71SS07I)	LACHESIS MUTA VENOM	8 [hp_X] in 59 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:63083-6022-2	59 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(63083-6022)