

# **PNEUMONIA/MENINGITIS NOSODE COMBINATION 9414- pneumonia/meningitis nosode 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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## **R414**

### **ACTIVE INGREDIENTS**

Aconitum napellus 3X  
Bryonia 3X  
Gelsemium sempervirens 3X  
Natrum carbonicum 3X  
Meningococcinum 12X  
Pneumococcinum 12X  
Silicea 12X

### **QUESTIONS**

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

### **INDICATIONS**

For the temporary relief of coughing, wheezing, chest congestion, occasional headache, fatigue, nausea, or vomiting.\*

\*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.

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### **DIRECTIONS**

Place drops under tongue 30 minutes before/after meals. Adults and children 12 years

and over: Take 10 to 15 drops once weekly or monthly. If mild symptoms are present, 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  
Pneumonia/Meningitis Nosode Combination  
Homeopathic Remedy  
1 FL. OZ. (29.5 mL)



PNEUMONIA/MENINGITIS NOSODE COMBINATION 9414			
pneumonia/meningitis nosode combination liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-9414
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
<b>ACONITUM NAPELLUS WHOLE</b> (UNII: U0NQ8555JD) (ACONITUM NAPELLUS WHOLE - UNII:U0NQ8555JD)	ACONITUM NAPELLUS WHOLE	3 [hp_X] in 29.5 mL
<b>BRYONIA ALBA ROOT</b> (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)	BRYONIA ALBA ROOT	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>SODIUM CARBONATE</b> (UNII: 45P3261C7T) (CARBONATE ION - UNII:7UJQ5OPE7D)	SODIUM CARBONATE	3 [hp_X] in 29.5 mL
<b>NEISSERIA MENINGITIDIS GROUP A CAPSULAR POLYSACCHARIDE ANTIGEN</b> (UNII: 1I86B47NY4) (NEISSERIA MENINGITIDIS GROUP A CAPSULAR POLYSACCHARIDE ANTIGEN - UNII:1I86B47NY4)	NEISSERIA MENINGITIDIS GROUP A CAPSULAR POLYSACCHARIDE ANTIGEN	12 [hp_X] in 29.5 mL
<b>STREPTOCOCCUS PYOGENES</b> (UNII: LJ2LP0YL98) (STREPTOCOCCUS PYOGENES - UNII:LJ2LP0YL98)	STREPTOCOCCUS PYOGENES	12 [hp_X] in 29.5 mL
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	12 [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:63083-9414-1	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(63083-9414)