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.

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
ACONITUM NAPELLUS WHOLE (UNII: U0NQ8555JD) (ACONITUM NAPELLUS WHOLE - UNII:U0NQ8555JD)	ACONITUM NAPELLUS WHOLE	3 [hp_X] in 29.5 mL
BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)	BRYONIA ALBA ROOT	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:7UJQ5OPE7D)	SODIUM CARBONATE	3 [hp_X] in 29.5 mL
NEISSERIA MENINGITIDIS GROUP A CAPSULAR POLYSACCHARIDE ANTIGEN (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
STREPTOCOCCUS PYOGENES (UNII: LJ2LP0YL98) (STREPTOCOCCUS PYOGENES - UNII:LJ2LP0YL98)	STREPTOCOCCUS PYOGENES	12 [hp_X] in 29.5 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	12 [hp_X] in 29.5 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (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)