POLLINOSIS COMBINATION 9261- pollinosis 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.

R261

ACTIVE INGREDIENTS

Pothos foetidus 5X
Arsenicum album 6X
Euphrasia officinalis 6X
Ignatia amara 6X
Kali phosphoricum 6X
Lycopodium clavatum 6X
Sanguinaria canadensis 6X
Sulphur 6X
Histaminum hydrochloricum 9X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of runny nose, nasal congestion, headache, or itchy and watery eyes due to hay fever or other upper respiratory allergies.*

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

WARNINGS

If symptoms do not improve or are accompanied by a fever, 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

40% ethanol, purified water.

LABEL

Est 1985

Professional Formulas

Complementary Health

Pollinosis Combination

Homeopathic Remedy

1 FL. OZ. (29.5 mL)

POLLINOSIS COMBINATION 9261

pollinosis combination liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63083-9261
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SYMPLOCARPUS FOETIDUS ROOT (UNII: R88254608W) (SYMPLOCARPUS FOETIDUS ROOT - UNII:R88254608W)	SYMPLOCARPUS FOETIDUS ROOT	5 [hp_X] in 29.5 mL
ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) - UNII:C96613F5AV)	ARSENIC TRIOXIDE	6 [hp_X] in 29.5 mL
EUPHRASIA STRICTA (UNII: C9642I91WL) (EUPHRASIA STRICTA - UNII:C9642I91WL)	EUPHRASIA STRICTA	6 [hp_X] in 29.5 mL
STRYCHNOS IGNATII SEED (UNII: 1NM3M2487K) (STRYCHNOS IGNATII SEED - UNII:1NM3M2487K)	STRYCHNOS IGNATII SEED	6 [hp_X] in 29.5 mL
POTASSIUM PHOSPHATE, UNSPECIFIED FORM (UNII: B7862WZ632) (POTASSIUM PHOSPHATE, UNSPECIFIED FORM - UNII:B7862WZ632)	POTASSIUM PHOSPHATE, UNSPECIFIED FORM	6 [hp_X] in 29.5 mL
LYCOPODIUM CLAVATUM SPORE (UNII: C88X29Y479) (LYCOPODIUM CLAVATUM SPORE - UNII:C88X29Y479)	LYCOPODIUM CLAVATUM SPORE	6 [hp_X] in 29.5 mL
SANGUINARIA CANADENSIS ROOT (UNII: N9288CD508) (SANGUINARIA CANADENSIS ROOT - UNII:N9288CD508)	SANGUINARIA CANADENSIS ROOT	6 [hp_X] in 29.5 mL
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	6 [hp_X] in 29.5 mL
HISTAMINE DIHYDROCHLORIDE (UNII: 3POA0Q644U) (HISTAMINE - UNII:820484N8I3)	HISTAMINE DIHYDROCHLORIDE	9 [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:63083-9261-1	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(63083-9261)