

ALLERGY NOSODE 4017- allergy nosode 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.

N17

ACTIVE INGREDIENTS

House dust mix 12X (15% by volume)
Mold mix 12X (15% by volume)
Grass mix 12X (10% by volume)
Tree mix 12X (10% by volume)
Weed mix 12X (10% by volume)
Kali bichromicum 30X (5% by volume)
Hepar sulphuris calcareum 12X (5% by volume)
Histaminum 12X (5% by volume)
Silicea 12X (5% by volume)
Euphorbium officinalis 6X (5% by volume)
Guaiaacum 6X (5% by volume)
Luffa operculata 6X (5% by volume)
Sinapsis nigra 3X (5% by volume)

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of runny nose, nasal congestion, headache, itchy or watery eyes, or mild facial or ear pain 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.

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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

20% ethanol, purified water.

LABEL

Est 1985

Professional Formulas

Complementary Health

Allergy Nosode

Homeopathic Remedy

2 FL. OZ. (59 mL)



ALLERGY NOSODE 4017

allergy nosode liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-4017
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOUSE DUST (UNII: EYO007VX98) (HOUSE DUST - UNII:EYO007VX98)	HOUSE DUST	12 [hp_X] in 59 mL
STACHYBOTRYS CHARTARUM (UNII: HJ4L70T1ZP) (STACHYBOTRYS CHARTARUM - UNII:HJ4L70T1ZP)	STACHYBOTRYS CHARTARUM	12 [hp_X] in 59 mL
MEDICAGO SATIVA WHOLE (UNII: DJO934BRBD) (MEDICAGO SATIVA WHOLE - UNII:DJO934BRBD)	MEDICAGO SATIVA WHOLE	12 [hp_X] in 59 mL
FRAXINUS LATIFOLIA WHOLE (UNII: 024W88F557) (FRAXINUS LATIFOLIA WHOLE - UNII:024W88F557)	FRAXINUS LATIFOLIA WHOLE	12 [hp_X] in 59 mL
TARAXACUM OFFICINALE (UNII: 39981FM375) (TARAXACUM OFFICINALE - UNII:39981FM375)	TARAXACUM OFFICINALE	12 [hp_X] in 59 mL
POTASSIUM DICHROMATE (UNII: T4423S18FM) (DICHROMATE ION - UNII:9LKY4BFN2V)	POTASSIUM DICHROMATE	30 [hp_X] in 59 mL
CALCIUM SULFIDE (UNII: 1MBW07J51Q) (CALCIUM SULFIDE - UNII:1MBW07J51Q)	CALCIUM SULFIDE	12 [hp_X] in 59 mL
HISTAMINE (UNII: 820484N8I3) (HISTAMINE - UNII:820484N8I3)	HISTAMINE	12 [hp_X] in 59 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	12 [hp_X] in 59 mL
GUAIAC (UNII: 03C8A0DFJ8) (GUAIAC - UNII:03C8A0DFJ8)	GUAIAC	6 [hp_X] in 59 mL
LUFFA OPERCULATA FRUIT (UNII: C4MO6809HU) (LUFFA OPERCULATA FRUIT - UNII:C4MO6809HU)	LUFFA OPERCULATA FRUIT	6 [hp_X] in 59 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:63083-4017-2	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(63083-4017)

Revised: 1/2026

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