

**PARASITES/AMOEBA/PROTOZOAN NOSODE COMBINATION 9402-
parasites/amoeba/protozoan 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.

R402

ACTIVE INGREDIENTS

Echinacea angustifolia 3X
Belladonna 4X
Eucalyptus globulus 4X
Eupatorium perfoliatum 6X
Amoeba 12X
Malaria 12X
Toxoplasmosis 12X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of occasional headache, mild abdominal pain or cramping, minor muscle ache or pain, fatigue, flatulence, nausea or vomiting, or diarrhea.*

*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, stiff neck, 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

Parasites/Amoeba/Protozoan Nosode Combination

Homeopathic Remedy

1 FL. OZ. (29.5 mL)



PARASITES/AMOEBEA/PROTOZOAN NOSODE COMBINATION 9402

parasites/amoeba/protozoan nosode combination liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-9402
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ECHINACEA ANGUSTIFOLIA WHOLE (UNII: VB06AV5US8) (ECHINACEA ANGUSTIFOLIA WHOLE - UNII:VB06AV5US8)	ECHINACEA ANGUSTIFOLIA WHOLE	3 [hp_X] in 29.5 mL
AMARYLLIS BELLADONNA WHOLE (UNII: 11B89M58UN) (AMARYLLIS BELLADONNA WHOLE - UNII:11B89M58UN)	AMARYLLIS BELLADONNA WHOLE	4 [hp_X] in 29.5 mL
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6) (EUCALYPTUS GLOBULUS LEAF - UNII:S546YLW6E6)	EUCALYPTUS GLOBULUS LEAF	4 [hp_X] in 29.5 mL
EUPATORIUM PERFOLIATUM FLOWERING TOP (UNII: 1W0775VX6E) (EUPATORIUM PERFOLIATUM FLOWERING TOP - UNII:1W0775VX6E)	EUPATORIUM PERFOLIATUM FLOWERING TOP	6 [hp_X] in 29.5 mL
AMOEBA PROTEUS (UNII: APR291GFYV) (AMOEBA PROTEUS - UNII:APR291GFYV)	AMOEBA PROTEUS	12 [hp_X] in 29.5 mL
ANOPHELES QUADRIMACULATUS (UNII: B4ULB6ARCO) (ANOPHELES QUADRIMACULATUS - UNII:B4ULB6ARCO)	ANOPHELES QUADRIMACULATUS	12 [hp_X] in 29.5 mL
TOXOPLASMA GONDII (UNII: BMV90JF469) (TOXOPLASMA GONDII - UNII:BMV90JF469)	TOXOPLASMA GONDII	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:63083-9402-1	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(63083-9402)