

NITROGEN COMBINATION 9310- nitrogen 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.

R310

ACTIVE INGREDIENTS

Cimicifuga racemosa 3X
Gelsemium sempervirens 3X
Natrium carbonicum 3X
Amitrole 12X
Atrazine 12X
Silicea 12X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of abdominal pain, nausea or vomiting, diarrhea, or skin, eye, or upper respiratory irritation.*

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

WARNINGS

Consult a doctor if condition worsens or if symptoms persist. 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 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
Nitrogen Combination
Homeopathic Remedy
1 FL. OZ. (29.5 mL)



NITROGEN COMBINATION 9310

nitrogen combination liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-9310
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
			2 [mg, X]

BLACK COHOSH (UNII: K73E24S6X9) (BLACK COHOSH - UNII:K73E24S6X9)	BLACK COHOSH	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
AMITROLE (UNII: ZF80H5GXUF) (AMITROLE - UNII:ZF80H5GXUF)	AMITROLE	12 [hp_X] in 29.5 mL
DESETHYL ATRAZINE (UNII: 07PV14BK6X) (DESETHYL ATRAZINE - UNII:07PV14BK6X)	DESETHYL ATRAZINE	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:63083-9310-1	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(63083-9310)