

GLYOXAL COMBINATION 9303- glyoxal 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.

R303

ACTIVE INGREDIENTS

Gelsemium sempervirens 3X

Natrum carbonicum 3X

Nux vomica 6X

Chelidonium majus 10X

Dimethyl phthalate 12X

Formalinum 12X

Glyoxal 12X

Sulphur 12X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of nasal congestion or irritation, minor sore throat, red or irritated skin or eyes, abdominal pain, or nausea.*

*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

Glyoxal Combination

Homeopathic Remedy

1 FL. OZ. (29.5 mL)



GLYOXAL COMBINATION 9303

glyoxal combination liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-9303
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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
STRYCHNOS NUX-VOMICA SEED (UNII: 269XH13919) (STRYCHNOS NUX-VOMICA SEED - UNII:269XH13919)	STRYCHNOS NUX-VOMICA SEED	6 [hp_X] in 29.5 mL
CHELIDONIUM MAJUS WHOLE (UNII: 7E889U5RNN) (CHELIDONIUM MAJUS WHOLE - UNII:7E889U5RNN)	CHELIDONIUM MAJUS WHOLE	10 [hp_X] in 29.5 mL
DIMETHYL PHTHALATE (UNII: 08X7F5UDJM) (DIMETHYL PHTHALATE - UNII:08X7F5UDJM)	DIMETHYL PHTHALATE	12 [hp_X] in 29.5 mL
FORMALDEHYDE SOLUTION (UNII: T0H3L6C7I5) (FORMALDEHYDE SOLUTION - UNII:T0H3L6C7I5)	FORMALDEHYDE SOLUTION	12 [hp_X] in 29.5 mL
(4-METHOXYPHENYL)GLYOXAL (UNII: 0IE1619UGN) ((4-METHOXYPHENYL)GLYOXAL - UNII:0IE1619UGN)	(4-METHOXYPHENYL)GLYOXAL	12 [hp_X] in 29.5 mL
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	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:63083-9303-1	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(63083-9303)