

BELLADONNA COMBINATION 9210- belladonna 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.

R210

ACTIVE INGREDIENTS

Belladonna 3X, 10X, 30X, 200X, 1000X
Echinacea angustifolia 3X, 10X, 30X, 200X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For the temporary relief of mild to moderate pain and inflammation, or fever.*

*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
Belladonna Combination
Homeopathic Remedy
1 FL. OZ. (29.5 mL)



BELLADONNA COMBINATION 9210

belladonna combination liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-9210
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
AMARYLLIS BELLADONNA WHOLE (UNII: 11B89M58UN) (AMARYLLIS BELLADONNA WHOLE - UNII:11B89M58UN)		AMARYLLIS BELLADONNA WHOLE	3 [hp_X] in 29.5 mL
ECHINACEA ANGUSTIFOLIA WHOLE (UNII: VB06AV5US8) (ECHINACEA ANGUSTIFOLIA WHOLE - UNII:VB06AV5US8)		ECHINACEA ANGUSTIFOLIA WHOLE	3 [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:63083-9210-1	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(63083-9210)

Revised: 1/2026

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