

## **TEETHING DROPS 2120- teething drops 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.*

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## **C120**

### **ACTIVE INGREDIENTS**

Eugenol 3X  
Ulmus fulva 3X  
Chamomilla 3X, 6X, 12X  
Mentha piperita 4X  
Clematis erecta 6X  
Colocynthis 6X  
Cuprum metallicum 6X  
Myrrha 6X  
Bryonia 12X, 30X  
Calcarea carbonica 12X, 30X

### **QUESTIONS**

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

### **INDICATIONS**

For temporary relief of pain or irritability associated with teething.\*

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

### **WARNINGS**

Consult a doctor if accompanied by fever and symptoms such as lethargy, lack of appetite, vomiting, or diarrhea. 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 for up to 6 weeks. For immediate onset of symptoms, take 10 to 15 drops every 15 minutes up to 3 hours. For less severe symptoms, take 10-15 drops hourly up to 8 hours. 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**

10% ethanol, purified water, 8% vegetable glycerin (from soy).

**LABEL**

Est 1985  
Professional Formulas  
Complementary Health  
Teething Drops  
Homeopathic Remedy  
2 FL. OZ. (59 mL)



TEETHING DROPS 2120			
teething drops liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63083-2120

Route of Administration		ORAL		
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
EUGENOL (UNII: 3T8H1794QW) (EUGENOL - UNII:3T8H1794QW)		EUGENOL	3 [hp_X] in 59 mL	
ULMUS RUBRA BARK (UNII: 91QY4PXU8Q) (ULMUS RUBRA BARK - UNII:91QY4PXU8Q)		ULMUS RUBRA BARK	3 [hp_X] in 59 mL	
ARTEMISIA ANNUA FLOWERING TOP (UNII: 0UQK6O82OW) (ARTEMISIA ANNUA FLOWERING TOP - UNII:0UQK6O82OW)		ARTEMISIA ANNUA FLOWERING TOP	3 [hp_X] in 59 mL	
MENTHA X PIPERITA WHOLE (UNII: 79M2M2UDA9) (MENTHA X PIPERITA WHOLE - UNII:79M2M2UDA9)		MENTHA X PIPERITA WHOLE	4 [hp_X] in 59 mL	
CLEMATIS RECTA FLOWERING TOP (UNII: 396421SP9F) (CLEMATIS RECTA FLOWERING TOP - UNII:396421SP9F)		CLEMATIS RECTA FLOWERING TOP	6 [hp_X] in 59 mL	
CITRULLUS COLOCYNTHIS FRUIT PULP (UNII: 23H32AOH17) (CITRULLUS COLOCYNTHIS FRUIT PULP - UNII:23H32AOH17)		CITRULLUS COLOCYNTHIS FRUIT PULP	6 [hp_X] in 59 mL	
COPPER (UNII: 789U1901C5) (COPPER - UNII:789U1901C5)		COPPER	6 [hp_X] in 59 mL	
MYRRH (UNII: JC71GJ1F3L) (MYRRH - UNII:JC71GJ1F3L)		MYRRH	6 [hp_X] in 59 mL	
BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)		BRYONIA ALBA ROOT	12 [hp_X] in 59 mL	
OYSTER SHELL CALCIUM CARBONATE, CRUDE (UNII: 2E32821G6I) (OYSTER SHELL CALCIUM CARBONATE, CRUDE - UNII:2E32821G6I)		OYSTER SHELL CALCIUM CARBONATE, CRUDE	12 [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:63083-2120-2	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(63083-2120)

Revised: 1/2026

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