

VERTIGO SYNCOPE DROPS 2115- vertigo syncope 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.

C115

ACTIVE INGREDIENTS

Arnica montana 3X
Hydrocotyle asiatica 3X
Phosphoricum acidum 6X
Ambra grisea 12X
Argentum nitricum 12X
Cocculus indicus 12X
Conium maculatum 12X
Lac caninum 12X
Theridion 12X

QUESTIONS

Professional Formulas

PO Box 2034 Lake Oswego, OR 97035

INDICATIONS

For temporary relief of occasional dizziness or lightheadedness, loss of balance, nausea, or vomiting.*

*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, persistent headache, stiff neck, or changes in vision or hearing, 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 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

20% ethanol, purified water.

LABEL

Est 1985

Professional Formulas

Complementary Health

Vertigo Syncope Drops

Homeopathic Remedy

2 FL. OZ. (59 mL)



VERTIGO SYNCOPE DROPS 2115

vertigo syncope drops liquid

Product Information

Product Type		HUMAN OTC DRUG	Item Code (Source)		NDC:63083-2115
Route of Administration		ORAL			
Active Ingredient/Active Moiety					
Ingredient Name			Basis of Strength	Strength	
ARNICA MONTANA WHOLE (UNII: O80TY208ZW) (ARNICA MONTANA WHOLE - UNII:O80TY208ZW)			ARNICA MONTANA WHOLE	3 [hp_X] in 59 mL	
CENTELLA ASIATICA LEAF (UNII: 6810070TYD) (CENTELLA ASIATICA LEAF - UNII:6810070TYD)			CENTELLA ASIATICA LEAF	3 [hp_X] in 59 mL	
PHOSPHORIC ACID (UNII: E4GA8884NN) (PHOSPHORIC ACID - UNII:E4GA8884NN)			PHOSPHORIC ACID	6 [hp_X] in 59 mL	
AMBERGRIS (UNII: XTC0D02P6C) (AMBERGRIS - UNII:XTC0D02P6C)			AMBERGRIS	12 [hp_X] in 59 mL	
SILVER NITRATE (UNII: 95IT3W8JZE) (SILVER CATION - UNII:57N7B0K90A)			SILVER NITRATE	12 [hp_X] in 59 mL	
ANAMIRTA COCCULUS FRUIT (UNII: 3E8XBL6YYK) (ANAMIRTA COCCULUS FRUIT - UNII:3E8XBL6YYK)			ANAMIRTA COCCULUS FRUIT	12 [hp_X] in 59 mL	
CONIUM MACULATUM FLOWERING TOP (UNII: Q28R5GF371) (CONIUM MACULATUM FLOWERING TOP - UNII:Q28R5GF371)			CONIUM MACULATUM FLOWERING TOP	12 [hp_X] in 59 mL	
CANIS LUPUS FAMILIARIS MILK (UNII: G39P120JQT) (CANIS LUPUS FAMILIARIS MILK - UNII:G39P120JQT)			CANIS LUPUS FAMILIARIS MILK	12 [hp_X] in 59 mL	
THERIDION CURASSAVICUM (UNII: 9Z8D3HEM8L) (THERIDION CURASSAVICUM - UNII:9Z8D3HEM8L)			THERIDION CURASSAVICUM	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:63083-2115-2	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(63083-2115)

Revised: 8/2019

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