

COCCULUS CONIUM SPECIAL ORDER- cocculus conium special order liquid
Uriel Pharmacy Inc.

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.

Cocculus Conium Special Order

Directions: FOR ORAL USE.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredients: Cocculus e fruct. 4X, Conium mac. 4X, Ambra grisea 7X, Petroleum 9X

Inactive Ingredients: Water, Salt

Use: Temporary relief of dizziness.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use.

Questions? Call 866.642.2858 Uriel, East Troy, WI 53120 www.urielpharmacy.com

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 Uriel, East Troy, WI 53120
www.urielpharmacy.com Lot:



**Cocculus
 Conium
 Special Order**

Homeopathic Ampules
 net vol. 0.3 fl. oz (10 x 1 ml)

Cocculus Conium s.o.

COCCULUS CONIUM SPECIAL ORDER

cocculus conium special order liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-3108
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ANAMIRTA COCCULUS WHOLE (UNII: 8O4P2U3QO2) (ANAMIRTA COCCULUS WHOLE - UNII:8O4P2U3QO2)	ANAMIRTA COCCULUS WHOLE	4 [hp_X] in 1 mL
CONIUM MACULATUM ROOT (UNII: TTE3RU7P5P) (CONIUM MACULATUM ROOT - UNII:TTE3RU7P5P)	CONIUM MACULATUM ROOT	4 [hp_X] in 1 mL

AMBERGRIS (UNII: XTC0D02P6C) (AMBERGRIS - UNII:XTC0D02P6C)	AMBERGRIS	9 [hp_X] in 1 mL
LIQUID PETROLEUM (UNII: 6ZAE7X688J) (LIQUID PETROLEUM - UNII:6ZAE7X688J)	LIQUID PETROLEUM	9 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-3108-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-3108)

Revised: 5/2018

Uriel Pharmacy Inc.