

CUPRUM ACETICUM NICOTIANA SPECIAL ORDER- cuprum aceticum nicotiana 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.

Cuprum Aceticum Nicotiana Special Order

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops.
Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredients: Cuprum aceticum 6X, Lobelia e pl. tota 6X, Renes 6X, Nicotiana e fol. 10X, Arsenicum album 12X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol

Use: Temporary relief of headache.

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. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858

Made with care by Uriel, East Troy, WI 53120

www.urielpharmacy.com

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-3214
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CUPRIC ACETATE (UNII: 39M11XPH03) (CUPRIC CATION - UNII:8CBV67279L)	CUPRIC ACETATE	6 [hp_X] in 1 mL
LOBELIA SPICATA LEAF (UNII: 1G4GK01F67) (LOBELIA SPICATA LEAF - UNII:1G4GK01F67)	LOBELIA SPICATA LEAF	6 [hp_X] in 1 mL
PORK KIDNEY (UNII: X7BCI5P86H) (PORK KIDNEY - UNII:X7BCI5P86H)	PORK KIDNEY	6 [hp_X] in 1 mL
TOBACCO LEAF (UNII: 6YR2608RSU) (TOBACCO LEAF - UNII:6YR2608RSU)	TOBACCO LEAF	10 [hp_X] in 1 mL
ARSENIC TRIOXIDE (UNII: S7V92P67HO) (ARSENIC CATION (3+) - UNII:C96613F5AV)	ARSENIC TRIOXIDE	12 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-3214-3	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	

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(48951-3214)

Revised: 5/2018

Uriel Pharmacy Inc.