

**CUPRUM ACETICUM ZINCUM VALERIANICUM- cuprum aceticum zincum valerianicum 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 Zincum valerianicum**

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 (Copper acetate) 5X, Zincum valerianicum (Valerate of zinc) 5X

Inactive Ingredients: Distilled water, Organic cane alcohol

Use: Temporary relief of cramps.

**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

Label:  
 Use: Temporary relief of cramps.  
 Inactive Ingredients: Distilled water, 15% Organic cane alcohol  
 Active Ingredients: Cuprum aceticum (Copper acetate) 5X, Zincum valerianicum (Valerate of zinc) 5X  
 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.



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cuprum aceticum zincum valerianicum liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:48951-3143
<b>Route of Administration</b>	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
CUPRIC ACETATE (UNII: 39M11XPH03) (CUPRIC CATION - UNII:8CBV67279L)			CUPRIC ACETATE	5 [hp_X] in 1 mL
ZINC VALERATE DIHYDRATE (UNII: MN0RX54EQA) (VALERIC ACID - UNII:GZK92PJM7B)			ZINC VALERATE DIHYDRATE	5 [hp_X] in 1 mL
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
ALCOHOL (UNII: 3K9958V90M)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-3143-3	60 mL in 1 BOTTLE, GLASS; 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(48951-3143)

Revised: 4/2018

Uriel Pharmacy Inc.