

**CINIS COMP.- cinis comp. 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.*

**Cinis comp.**

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: Cinis Equiseti (Ash of horsetail herb) 6X, Cinis Quercus (Decoction of oak bark ash) 6X, Cinis Tabaci (Tobacco ash) 6X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol, Lactose

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. Contains traces of lactose. 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 Lot:



<b>CINIS COMP.</b>			
cinis comp. liquid			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:48951-3242
<b>Route of Administration</b>	ORAL		
<b>Active Ingredient/Active Moiety</b>			
Ingredient Name		Basis of Strength	Strength

<b>EQUISETUM ARVENSE TOP</b> (UNII: 1DP6Y6B65Z) (EQUISETUM ARVENSE TOP - UNII:1DP6Y6B65Z)	EQUISETUM ARVENSE TOP	6 [hp_X] in 1 mL
<b>QUERCUS ROBUR WHOLE</b> (UNII: R7QMG0BT2W) (QUERCUS ROBUR WHOLE - UNII:R7QMG0BT2W)	QUERCUS ROBUR WHOLE	6 [hp_X] in 1 mL
<b>TOBACCO LEAF</b> (UNII: 6YR2608RSU) (TOBACCO LEAF - UNII:6YR2608RSU)	TOBACCO LEAF	6 [hp_X] in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>LACTOSE, UNSPECIFIED FORM</b> (UNII: J2B2A4N98G)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-3242-3	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(48951-3242)

Revised: 3/2021

Uriel Pharmacy Inc.