

**CINNAMIC ACID- cinnamic acid liquid**  
**Deseret Biologicals, 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.*

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**DRUG FACTS:**

**ACTIVE INGREDIENTS:**

Cinnamic Acid 6X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

**HOMEOPATHIC INDICATIONS:**

For temporary relief of skin rashes, general fatigue, craving for fruit, headaches, and indigestion.\*\*

\*\*These statements are based upon traditional homeopathic practice. They have not been reviewed by the Food and Drug Administration.

**WARNINGS:**

**Keep out of reach of children.** In case of overdose, contact a physician or Poison Control Center right away.

**If pregnant or breast feeding,** ask a health professional before use.

Tamper seal: "Sealed for Your Protection."

Do not use if seal is broken or missing.

**KEEP OUT OF REACH OF CHILDREN:**

**Keep out of reach of children.** In case of overdose, contact a physician or Poison Control Center right away.

**DIRECTIONS:**

1-10 drops under the tongue, 3 times a day or as directed by a health professional. Consult a physician for use in children under 12 years of age.

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**INACTIVE INGREDIENTS:**

Demineralized Water, 25% Ethanol

**QUESTIONS:**

Dist. By: Deseret Biologicals, Inc.  
469 W. Parkland Drive  
Sandy, UT 84070 [www.desbio.com](http://www.desbio.com)

**PACKAGE DISPLAY LABEL:**

**DESBIO**

NDC 43742-0659-1

HOMEOPATHIC

**CINNAMIC ACID**

1 FL OZ (30 ml)

**WARNINGS:**

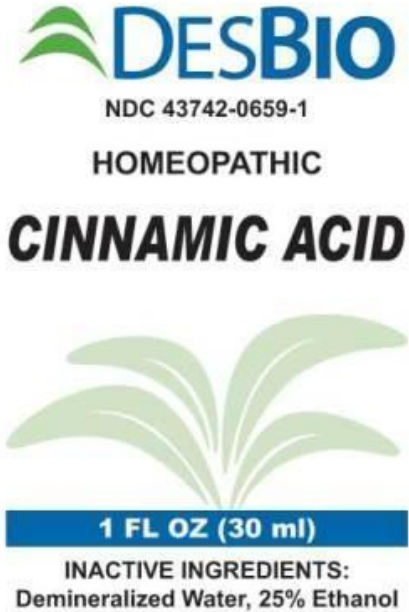
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**CINNAMIC ACID**

cinnamic acid liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:43742-0659
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
CINNAMIC ACID (UNII: U14A832J8D) (CINNAMIC ACID - UNII:U14A832J8D)	CINNAMIC ACID	6 [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:43742-0659-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	11/11/2015	11/15/2022

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		11/11/2015	11/15/2022

**Labeler** - Deseret Biologicals, Inc. (940741853)

**Registrant** - Apotheca Company (844330915)

### Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0659) , api manufacture(43742-0659) , label(43742-0659) , pack(43742-0659)

Revised: 11/2018

Deseret Biologicals, Inc.