# FORMALDEHYDE- formalinum 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:**

Formalinum 6X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

#### **INDICATIONS:**

For temporary relief of reactions related to formaldehyde sensitivity including burning eyes, crying, dizziness, muscle spasms, shakiness and throat inflammation.\*\*

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

#### **WARNINGS:**

**Keep out of reach of children.** In case of overdose, contact physician or a 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 physician or a 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

#### **PACKAGE LABEL DISPLAY:**

**DESBIO** 

NDC 43742-0774-1

**HOMEOPATHIC** 

**FORMALDEHYDE** 

1 FL OZ (30 ml)

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#### LOT:

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#### **FORMALDEHYDE**

Droduct Information

formalinum liquid

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-0774

Route of Administration ORAL

#### **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
FORMALDEHYDE (UNII: 1HG84L3525) (FORMALDEHYDE - UNII:1HG84L3525)	FORMALDEHYDE	6 [hp_X] in 1 mL

	Inactive Ingredients		
ı	Ingredient Name	Strength	
ı	WATER (UNII: 059QF0KO0R)		
ı	ALCOHOL (UNII: 3K9958V90M)		

	Packaging				
# Item Code		Package Description	Marketing Start Date	Marketing End Date	
	NDC:43742- 0774-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	04/14/2017	11/0 1/20 22	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/22/2017	11/0 1/20 22

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

### **Registrant** - Apotheca Company (844330915)

Establishment			
Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0774), api manufacture(43742-0774), label(43742-0774), pack(43742-0774)

Revised: 9/2018 Deseret Biologicals, Inc.