CONIFERYL ALCOHOL- coniferyl alcohol 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.

Drug Facts:

ACTIVE INGREDIENTS:

Coniferyl Alcohol 6X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

INDICATIONS:

For temporary relief of symptoms caused by allergies including rhinitis, congestion, headaches, digestive problems and rashes.

WARNINGS:

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

If pregnant or breast-feeding, seek advice of a health professional before use.

Tamper seal: "Sealed for Your Protection." Do not use if seal is broken or missing.

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.

INACTIVE INGREDIENTS:

Demineralized Water, 25% Ethanol.

KEEP OUT OF REACH OF CHILDREN:

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

INDICATIONS:

For temporary relief of symptoms caused by allergies including rhinitis, congestion, headaches, digestive problems and rashes.

QUESTIONS:

Dist. By: Deseret Biologicals, Inc.

469 W. Parkland Drive

Sandy, UT 84070 www.desbio.com

PACKAGE LEBEL DISPLAY:

DES**BIO**

NDC 43742-0185-1

HOMEOPATHIC

CONIFERYL

ALCOHOL

1 FL OZ (30 ml)

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CONIFERYL ALCOHOL
coniferyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43742-0185
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CONIFERYL ALCOHOL (UNII: E7SM92591P) (CONIFERYL ALCOHOL - UNII:E7SM92591P)	CONIFERYL ALCOHOL	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:43742-0185-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	10/29/2012	11/22/2021

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/29/2012	11/22/2021

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company (844330915)

Establishment

Name	Address	ID/FEI	Business Operations
Apotheca Company		844330915	manufacture(43742-0185) , api manufacture(43742-0185) , label(43742-0185) , pack(43742-0185)

Revised: 12/2018 Deseret Biologicals, Inc.