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:

DESBIO

NDC 43742-0185-1

HOMEOPATHIC

CONIFERYL

ALCOHOL

1 FL OZ (30 ml)

WARNINGS:

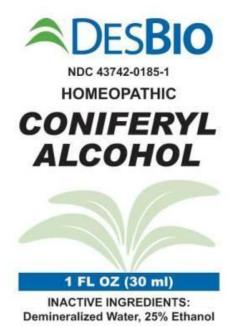
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LOT:

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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 CONIFERYL ALCOHOL (UNII: E7SM92591P) (CONIFERYL ALCOHOL UNII:E7SM92591P) CONIFERYL ALCOHOL | CONIFERYL | 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 BOTTL Product		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	10/29/2012	11/22/2021			
Marketing Information							
	Marketing Categ	ory Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			

10/29/2012

11/22/2021

Labeler - Deseret Biologicals, Inc. (940741853)

unapproved homeopathic

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.