**CONIFERYL ALCOHOL PHENOLIC- coniferyl alcohol liquid**
**Energique, 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 INGREDIENT:

**(in each drop):** 24.98% of Coniferyl Alcohol 12X, 30X, 12C, 30C; 0.10% of Coniferyl Alcohol 6X.

## INDICATIONS:

May temporarily relieve symptoms associated with coniferyl alcohol reactions.**
**Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

## WARNINGS:

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

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

Do not use if tamper evident seal is broken or missing. Store in a cool, dry place.

## KEEP OUT OF REACH OF CHILDREN:

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

## DIRECTIONS:

Adults and children 5 to 10 drops orally, 1 time daily or as otherwise directed by a health care professional. If symptoms persist for more than 7 days, consult your health care professional. Consult a physician for use in children under 12 years of age.

## INDICATIONS:

May temporarily relieve symptoms associated with coniferyl alcohol reactions.**
**Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

## INACTIVE INGREDIENTS:

Demineralized water, 20% Ethanol

**QUESTIONS:**

Dist. by Energique, Inc.

201 Apple Blvd

Woodbine, IA 51579 **800-869-8078**

**PACKAGE LABEL DISPLAY:**

**ENERGIQUE**

SINCE 1987

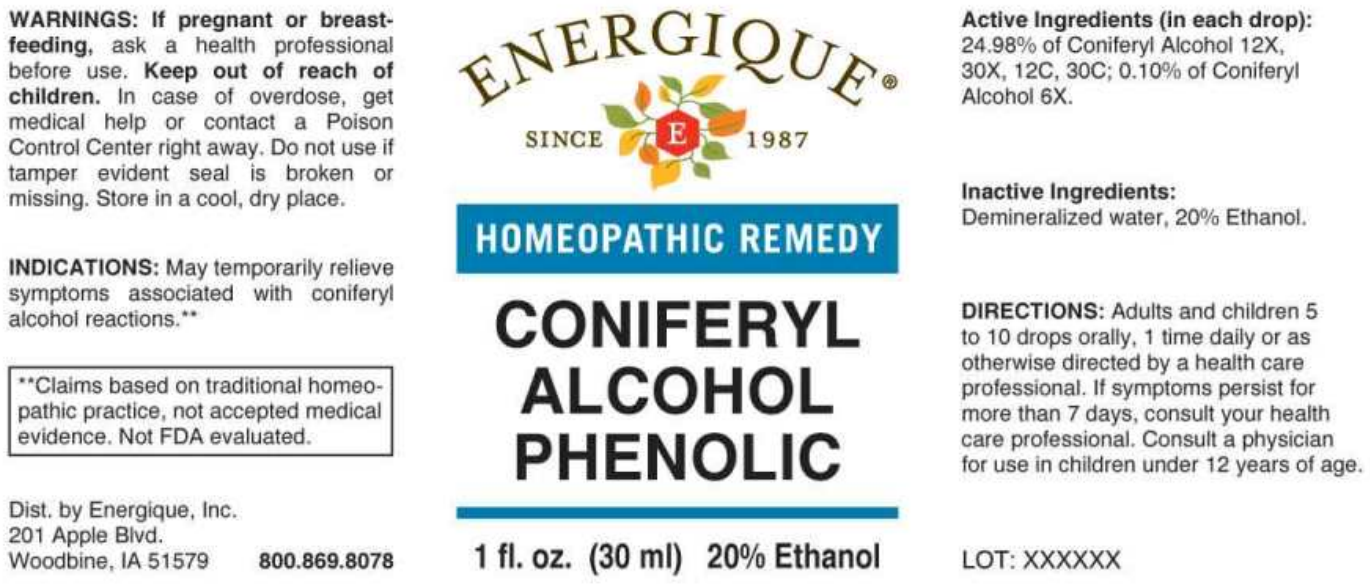
**HOMEOPATHIC REMEDY**

**CONIFERYL**

**ALCOHOL**

**PHENOLIC**

**1 fl. oz. (30 ml)**




## CONIFERYL ALCOHOL PHENOLIC

coniferyl alcohol liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:44911-0253 |
| **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:44911-0253-1 | 30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product | 08/31/2015 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved homeopathic | | 08/31/2015 | |

## Labeler - Energique, Inc. (789886132)

## Registrant - Apotheca Company (844330915)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Apotheca Company | | 844330915 | manufacture(44911-0253) , api manufacture(44911-0253) , label(44911-0253) , pack(44911-0253) |

Revised: 10/2024 Energique, Inc.