

FAIRY HANDS HAND SANITIZER- alcohol liquid ECBLEND LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Hand Sanitizer

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Alcohol 80% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

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Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel

00 mL NDC: 00000-000-00

74296-101-03 1 BOTTLE 120mL

74296-101-02 1 BOTTLE 60mL

74296-101-07 1 BOTTLE 3785mL

74296-101-04 1 BOTTLE 250mL

74296-101-05 1 BOTTLE 500mL

74296-101-06 1 BOTTLE 1000mL

Drug Facts

Active ingredient[s]	Purpose
Alcohol 80% v/v.....	Antiseptic

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Inactive Ingredients glycerin, hydrogen peroxide, purified water USP

Questions? (541) 393-9840

Manufactured & Distributed by
ECBlend LLC 226 N Ross Ln, Medford, OR 97501
ECBlend.com

FairyHands



HAND Sanitizer

**ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION**
Non-sterile Solution

EXP:

BATCH ID

2 FL OZ (60 ML)
NDC 74296-101-02



Drug Facts

Active ingredient[s]	Purpose
Alcohol 80% v/v.....	Antiseptic

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HAND Sanitizer

ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION
Non-sterile Solution

EXP:

BATCH ID

34 FL OZ (1000 ML)
NDC 74296-101-06



Drug Facts

Active ingredient[s]	Purpose
Alcohol 80% v/v.....	Antiseptic

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HAND Sanitizer

ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION
Non-sterile Solution

EXP:

BATCH ID

8.45 FL OZ (250 ML)
NDC 74296-101-04



Drug Facts

Active ingredient[s]	Purpose
Alcohol 80% v/v.....	Antiseptic

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Inactive Ingredients glycerin, hydrogen peroxide, purified water USP

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HAND Sanitizer

ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION
Non-sterile Solution

EXP:

BATCH ID

ONE GALLON (3.78L)
NDC 74296-101-07



Drug Facts

Active ingredient[s]	Purpose
Alcohol 80% v/v.....	Antiseptic

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- Store between 15-30C (59-86F)
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Inactive Ingredients glycerin, hydrogen peroxide, purified water USP

Questions? (541) 393-9840

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HAND Sanitizer

ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION
Non-sterile Solution

EXP:

BATCH ID

16.9 FL OZ (500 ML)
NDC 74296-101-05



Drug Facts

Active ingredient[s] *Purpose*
Alcohol 80% v/v.....Antiseptic

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Warnings
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FAIRY HANDS HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74296-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74296-101-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:74296-101-03	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
3	NDC:74296-101-04	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
4	NDC:74296-101-05	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
5	NDC:74296-101-06	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
6	NDC:74296-101-07	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - ECBLEND LLC (063972602)

Establishment

Name	Address	ID/FEI	Business Operations
ECBLEND LLC		063972602	manufacture(74296-101)

Revised: 3/2020

ECBLEND LLC