

HAND SANITIZER- alcohol liquid
Cnt Dream. Co., Ltd

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.

LOVLUV Hand Sanitizer 3.4 FL.OZ. (100 ml)

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

HAND SANITIZER

Non-Sterile Solution
Helps Reduce Bacteria

Moisturizing
+ Non-Sticky

LOVLUV

Net 3.4 FLOZ. (100 mL)

Drug Facts

Active Ingredient

Ethyl Alcohol 80% v/v

Uses

Hand sanitizer to help reduce cause disease. For use when s

Warnings

For external use only. Fla
Keep away from heat or fl

Do not use

- In children less than 2 mo
- On open skin wounds

When using this product l
mouth. In case of contact v
thoroughly with water.

Stop use and ask a doctor

These may be signs of a se
Keep out of reach of child
help or contact a Poison Cc

Directions

- Place enough product on
- Rub hands together until dr
- Supervise children under
- this product to avoid swallo

Other Information

- Store between 15-30°C (!
- Avoid freezing and excess

Inactive Ingredients W

Carbomer, Aminomethyl Pro
Aloe Barbadensis Leaf Extr
Extract, Rosmarinus Officina

Questions or Commer

Visit www.lovluvus.com

Distributed by CGETC INC.
City of Industry, CA

Made In Korea NDC No. 71909-0017

Purpose

Antiseptic, Hand Sanitizer

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Non-Sterile Solution
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Net 3.4 FLOZ. (100 ml)

Drug Facts

Active Ingredient

Ethyl Alcohol 80% v/v

Uses

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 old
- On open skin wounds

When using this product in the mouth. In case of contact with eyes, flush thoroughly with water.

Stop use and ask a doctor

These may be signs of a severe allergic reaction. Keep out of reach of children. Get medical help or contact a Poison Control Center if you experience these symptoms.

Directions

- Place enough product on palm of one hand to cover both hands.
- Rub hands together until dry.
- Supervise children under 12 years old when using this product to avoid swallowing.

Other Information

- Store between 15-30°C (59-86°F).
- Avoid freezing and excessive heat.

Inactive Ingredients

Water, Carbomer, Aminomethyl Propane Carboxylic Acid, Aloe Barbadensis Leaf Extract, Rosmarinus Officinalis Extract, Fragrance.

Questions or Comments

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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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Do not use

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- On open skin wounds

When using this product in the mouth, in case of contact with eyes, rinse 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. For more information, call 1-800-LOVLUV or visit www.lovluvus.com.

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Directions

- Place enough product on palm of one hand to cover both hands.
- Rub hands together until dry.
- Supervise children under 12 years of age when using this product to avoid swallowing.

Other Information

- Store between 15-30°C (59-86°F).
- Avoid freezing and excessive heat.

Inactive Ingredients

Water, Carbomer, Aminomethyl Propane Carboxylic Acid, Aloe Barbadensis Leaf Extract, Rosmarinus Officinalis Extract, Glycerin, Ethyl Alcohol, Fragrance, Potassium Sorbate, Sodium Benzoate, Sodium Chloride, Sodium Hydroxide, Tetrasodium EDTA, Xanthan Gum.

Questions or Comments

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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.

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Other information

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

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Inactive ingredients

Water, Glycerin, Fragrance, Carbomer, Aminomethyl Propanol, Sodium Hyaluronate, Aloe Barbadensis Leaf Extract, Camellia Sinensis Leaf Extract, Rosmarinus Officinalis (Rosemary) Leaf Extract.

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Package Label - Principal Display Panel

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Made In Korea NDC No. 71909-0017

HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71909-0017
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
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71909-0017-3	48 in 1 CASE	03/30/2020	
1	NDC:71909-0017-2	12 in 1 CASE		
1	NDC:71909-0017-1	100 mL in 1 BOTTLE; Type 0: Not a Combination Product		

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Stop use and ask a doctor
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Directions

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Other Information

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Inactive Ingredients Water,
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Marketing Information

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

Labeler - Cnt Dream. Co., Ltd (694699750)

Registrant - Cnt Dream. Co., Ltd (694699750)

Establishment

Name	Address	ID/FEI	Business Operations
Cnt Dream. Co., Ltd		694699750	manufacture(71909-0017)

Revised: 4/2020

Cnt Dream. Co., Ltd