

LOVLUV INSTANT 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 Instant Hand Sanitizer 3.4 FL.OZ. (100 mL)

Active Ingredient(s)

Drug Facts	
Active Ingredient	Purpose
Ethyl Alcohol 80% v/v	Antiseptic
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 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.	
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 15-30°C (59-86°F) - Avoid freezing and excessive heat above 40°C (104°F)	
Inactive Ingredients Water, Glycerin, Fragrance, Carbomer, Aminomethyl Propanol, Sodium Hyaluronate, Aloe Barbadosis Leaf Extract, Camellia Sinensis Leaf Extract, Rosmarinus Officinalis (Rosemary) Leaf Extract.	
Questions or Comments? Call +82-41-520-8943 & Visit www.lovluvus.com	

Manufactured by
CNT Dream Co., Ltd
80 Baekseokgongdan 3-gil, Seobuk-gu,
Cheonan-si, Chungcheongnam-do,
Korea

Made in Korea

8 809605 874144

Alcohol 80% v/v. Purpose: Antiseptic

Purpose

LOVLUV

INSTANT

HAND SANITIZER

*Non-Sterile Solution
Help Reducing Bacteria*

**Moisturizing
Non-Sticky**

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Antiseptic, Hand Sanitizer

Use

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Package Label - Principal Display Panel

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Moisturizing
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LOVLUV INSTANT HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71909-4144
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	24.066 mL in 100 mL

Product Characteristics

Color		Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71909-4144-3	48 in 1 CASE	04/01/2020	
1	NDC:71909-4144-1	100 mL in 1 BOTTLE; Type 0: Not a Combination Product		

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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2020	

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

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

Establishment

Name	Address	ID/FEI	Business Operations
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