BEWELL HAND SANITIZER- is opropyl alcohol solution L.C. INDUSTRIES, INC.

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.

Active Ingredient(s)

Isopropyl Alcohol 75% v/v.

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, water





bewell HAND **SANITIZER**

- · Isopropyl Alcohol Antiseptic 75% Topical Solution
- · Non-Sterile Solution

2 FL. OZ. (59 mL)

Drug Facts

Active ingredient Isopropyl alcohol 75% v/v

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BEWELL HAND SANITIZER

isopropyl alcohol solution

Product Information

HUMAN OTC DRUG Product Type Item Code (Source) NDC:77477-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Basis of Strength Ingredient Name Strength ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL -ISOPROPYL 75 mL UNII:ND2M416302) ALCOHOL in 100 mL

Inactive Ingredients

Ingredient Name	Strength	
GLYCERIN (UNII: PDC6A3C0OX)		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)		
WATER (UNII: 059QF0KO0R)		

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:77477-001-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020		
Marketing Information				
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not fi	nal part333A	05/08/2020		

Labeler - L.C. INDUSTRIES, INC. (069952018)

Revised: 5/2020 L.C. INDUSTRIES, INC.