HAND SANITIZER- is opropyl alcohol liquid Brewer Science, 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.

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. Isopropyl Alcohol (75%, 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)

Isopropyl Alcohol 75% 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

3800 ml NDC: 78175-111-38

Isopropyl Alcohol Antiseptic 75% Topical Solution

Antiseptic Hand Rub Non-sterile Solution

3800 ml



2401 Brewer Drive - Rolla, MO 65401 Tel: 573-364-0300

TOPICAL

Drug Facts	
Active ingredient[s] Isopropyl alcohol 75% v/v	Purpose Antiseptic
Use[s]	
Health care personnel hand rub to help reduce bacteria that potentially can cause disease.	
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 15-30C (59-86F)	
Avoid freezing and excessive heat above 40C (104F)	
Inactive ingredients glycerin, hydrogen peroxide, purified water USP	

HAND SANITIZER

isopropyl alcohol liquid

Draduct Information

Route of Administration

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78175-112	

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
		2855.7 mL	
UNII:ND2M416302)	ALCOHOL	in 3800 mL	

Inactive Ingredients			
Ingredient Name	Strength		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	4.7538 mL in 3800 mL		
WATER (UNII: 059QF0KO0R)	730.74 mL in 3800 mL		
GLYCERIN (UNII: PDC6A3C0OX)	53.998 mL in 3800 mL		

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date

1 NDC:78175-112-38 3800 mL in 1 BOTTLE; Type 0: Not a Combination Product 07/15/2020

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/15/2020	

Labeler - Brewer Science, Inc. (019689330)

Registrant - Brewer Science, Inc. (019689330)

Establishment				
Name	Address	ID/FEI	Business Operations	
Brewer Science, Inc.		019689330	manufacture (78 175-112)	

Revised: 11/2020 Brewer Science, Inc.