HAND SANITIZER- alcohol liquid Kevin Sathre

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.

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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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

Package Label - Principal Display Panel



60 mL NDC: 55909-080-60

HAND SANITIZER alcohol liquid **Product Information** HUMAN OTC DRUG NDC:55909-080 **Product Type** Item Code (Source) **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			
Н	YDRO GEN PERO XID	E (UNII: BBX060AN9V)		0.125 mL in 100 mL				
W	ATER (UNII: 059QF0	KO0R)						
P #	ackaging Item Code	Package Description		eting Start	Marketing End			
1		60 mL in 1 BOTTLE; Type 1: Convenience Kit of Co- Package	03/30/202	Date 20	Date			
Marketing Information								
	Marketing Categor	y Application Number or Monograph Citation	Marketin	g Start Date	Marketing End Date			
0	TC monograph not fin	al part333A	03/30/2020					

Labeler - Kevin Sathre (024321536)

Establishment

Name	Address	ID/FEI	Business Operations
Sapthre Maple Syrup		024321536	manufacture(55909-080)

Revised: 4/2020

Kevin Sathre