HAND SANITIZER- alcohol liquid SMK Holdings LLC.

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: 77177-001-01



120 mL NDC: 77177-001-02



3785 mL NDC: 77177-001-03

	NEWI	Drug Facts Active Ingredientie Purpese Alcohol 80% v/vAntiseptic
-		Use(s)
		Hand sanitizer to help reduce bacteria that potentially can cause disease.
		For use when soap and water are not available.
		_ Warnings
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		De not use • In children less than 2 months of age
		On open skin wounds
	Hand Sanitizer	When using this product
		 Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.
· · · ·	Alcohol Antiseptic	Step ase and ask a doctor
	Topical Solution	 If irritation or rash occurs. These may be signs of a serious condition.
	Non-Sterile Solution	 If swallowed, get medical help or contact a Poison Control Center right away.
750		KEEP OUT OF REACH OF CHILDREN.
	TOPO O ONL	Directions
Park Plan		 Place enough product on hands to cover all surfaces. Rub hands together until dry.
y: Sugar Creek Ct. Indianapolia, i		 Supervise children under 6 years of age when using this product to avoid swallowing.
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		 Store between 15 - 30C (59 - 86F)
		 Avoid freezing and excessive heat above 40C (104F)
	LIQUID SANITIZER	Inactive Ingredients
	1 Gallon (128 fl oz)	glycerin, hydrogen percodde, purified water USP

alcohol liquid

Product Information								
Product Type		HUMAN OTC DRUG	Item Code (Source)		NDC:77177-001(NDC:74221-001)			
Route of Administration		TOPICAL						
Active Ingredient/Act	ive Moi	ety						
Ingredient Name			Basis	of Strength	Strength			
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL		80 mL in 100 mL			
Inactive Ingredients	Inactive Ingredients							
Ingredient Name				Strength				
GLYCERIN (UNII: PDC6A3C0OX)			1.45 mL in 100 mL					
HYDROGEN PEROXIDE (UNII: BBX060AN9V)				0.125 mL in 100 mL				
WATER (UNII: 059QF0KO0F	R)							
Packaging								
# Item Code		Package Description	Μ	larketin	g Start Date	Marketing End Date		

Marketing Catego OTC monograph not fin	al part333A	03/30/2020			
Marketing Catego					
Marketing Catego	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
Marketing Information					
3 NDC:77177-001-03	3785 mL in 1 JUG; Type 0: Not a Combination Product	03/30/2020			
2 NDC:77177-001-02	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020			
I NDC.//1//-001-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020			

Labeler - SMK Holdings LLC. (095712546)

Establishment

Name	Address	ID/FEI	Business Operations
SMK Holdings LLC		095712546	repack(77177-001)

Revised: 5/2020

SMK Holdings LLC.