HAND SANITIZER- alcohol gel Marquis Extraction Technology 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



Alcohol Antiseptic 80% **Topical Solution**

Antiseptic Hand Rub Non-sterile Solution 473 ml

Drug Facts						
Active ingredient[s]	Purpose					
Alcohol 80% v/v	Antiseptic					
Use[s] Health care personnel hand rub to help reduce bacteria that p disease	potentially can cause					
<i>Warnings</i> For external use only. Flammable. Keep away from heat or fla	ame					
Do not usein children less than 2 months of ageon 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						
Batch# 001 expires 03/07/22 Manufactured by: Marguis Inc.						

In the event of an adverse reaction, call: 888-925-7311

11953 Prairie Industrial Pkwy Hennepin, IL 61327



473 mL NDC: 76585-001-01

н	AND SANITI	ZER					
	ohol gel						
Р	roduct Informa	tion					
			HUMAN OTC DRUG	em Code	e (Source)	NDC:76585-001
Product Type Route of Administration			TOPICAL	Item Code (Source)			
1	ute of Auministre	luon					
A	ctive Ingredien	t/Active Moi	ety				
Ingredient Name				Basis	of Strength	Strength	
							80 mL in 100 mL
Inactive Ingredients							
		Ing	gredient Name				Strength
	L YCERIN (UNII: PDC					1.45 mL in 10	
	Y DROGEN PEROXI ATER (UNII: 059QF0		0 AN9 V)			0.125 mL in 1	00 mL
P	ackaging						
#	Item Code		Package Description			ting Start Date	Marketing End Dat
1	NDC:76585-001- 07	3785 mL in 1 BO	TTLE; Type 0: Not a Combination	Product	03/30/202	0	
2	NDC:76585-001-01	473 mL in 1 BOT	TLE; Type 0: Not a Combination Pr	oduct	03/30/2020		
3	NDC:76585-001- 03	236 mL in 1 BOT	TLE; Type 0: Not a Combination P	roduct	03/30/2020		
4	NDC:76585-001- 04	946 mL in 1 BO	TTLE; Type 0: Not a Combination P	roduct	03/30/2020		
	NDC:76585-001- 05	121133 mL in 1 B Product	OTTLE; Type 0: Not a Combination	1	03/30/2020		
	NDC:76585-001- 02	1040875 mL in 1	TANK; Type 0: Not a Combination	Product	03/30/2020		
'	NDC:76585-001- 08		TLE; Type 0: Not a Combination Pr		03/30/2020		
8	NDC:76585-001- 06	18927 mL in 1 B Product	OTTLE; Type 0: Not a Combination		03/30/2020		
9	NDC:76585-001- 09	500 mL in 1 BOT	TLE; Type 0: Not a Combination P	roduct	03/30/202	0	
Marketing Information							
-	Marketing Catego	ry Applicat	ion Number or Monograph Cita	tion 1	Marketin	g Start Date	Marketing End Date
0	ГС monograph not fi	nal part333A		0	3/30/2020		

Labeler - Marquis Extraction Technology LLC (117496233)

Establishment								
Name	Address	ID/FEI	Business Operations					
Marquis Extraction Technology LLC		117496233	manufacture(76585-001)					

Revised: 5/2020

Marquis Extraction Technology LLC