# HAND SANITIZER LIQUID- alcohol liquid Wm Refractories, S. de R.L.de C.V.

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.

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### **Hand Sanitizer Liquid**

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:

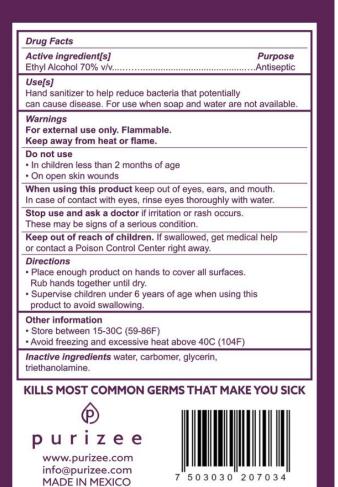
- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water 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)



20.29 fl.oz. Establishment approved by the FDA



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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#### **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.

approved by

the FDA

#### 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



**Drug Facts** Active ingredient[s] Ethyl Alcohol 70% v/v. Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available 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. · 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 water, carbomer, glycerin, KILLS MOST COMMON GERMS THAT MAKE YOU SICK purizee www.purizee.com info@purizee.com MADE IN MEXICO

250 mL NDC: 75260-322-08 330 mL NDC: 75260-322-09

600 ml

500 mL NDC: 75260-322-10 750 mL NDC: 75260-322-11 1893 mL NDC: 75260-322-12 3875 mL NDC: 75260-322-13 600 mL NDC: 75260-322-14

# HAND SANITIZER LIQUID

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75260-322
Poute 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		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL		
WATER (UNII: 059QF0KO0R)			

Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:75260-322-08	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:75260-322-09	330 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
3	NDC:75260-322-10	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
4	NDC:75260-322-11	750 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
5	NDC:75260-322-12	1893 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
6	NDC:75260-322-13	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
7	NDC:75260-322-14	600 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	03/30/2020		

# Labeler - Wm Refractories, S. de R.L.de C.V. (812611254)

# Registrant - Wm Refractories, S. de R.L.deC.V. (812611254)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
Wm Refractories, S. de R.L.deC.V.		8 126 11254	manufacture(75260-322)

Revised: 5/2020 Wm Refractories, S. de R.L.de C.V.