

**HAND SANITIZER- alcohol liquid**  
**Emergency Medical Solutions 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.*

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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.

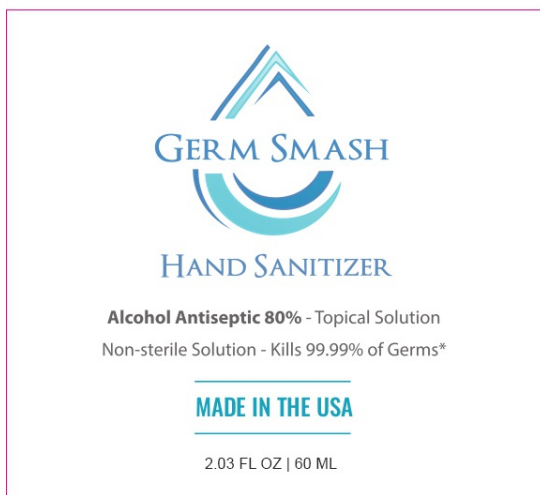
## Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

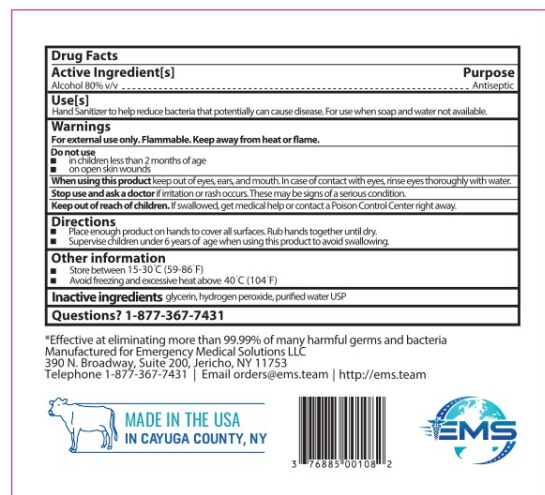
## Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

## Package Label - Principal Display Panel



FRONT

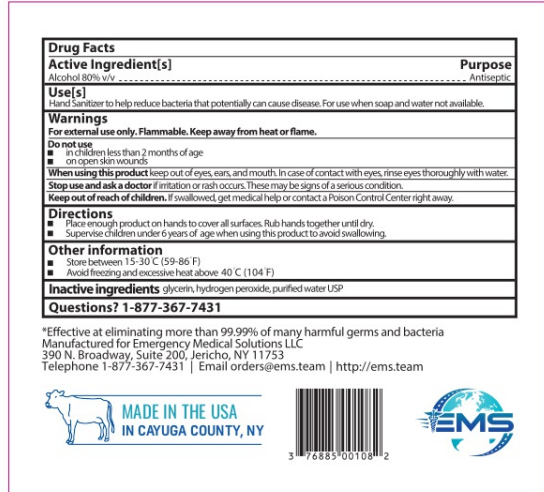


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LABEL BACKGROUNDS ARE 100% WHITE

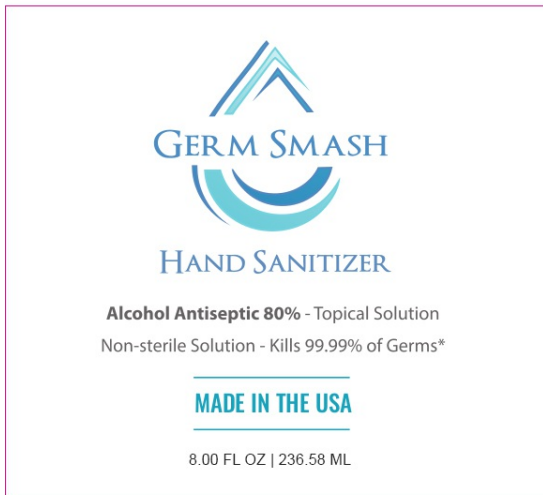


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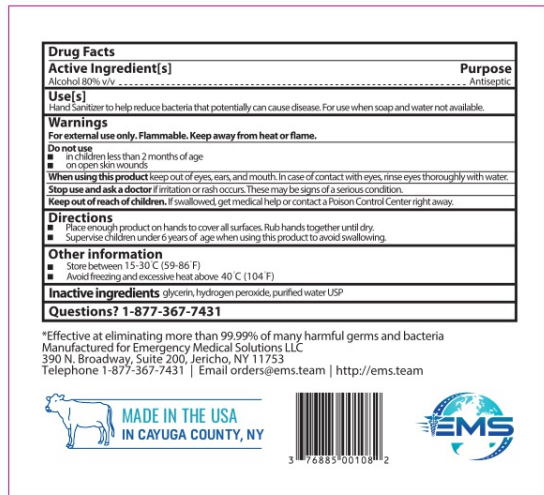


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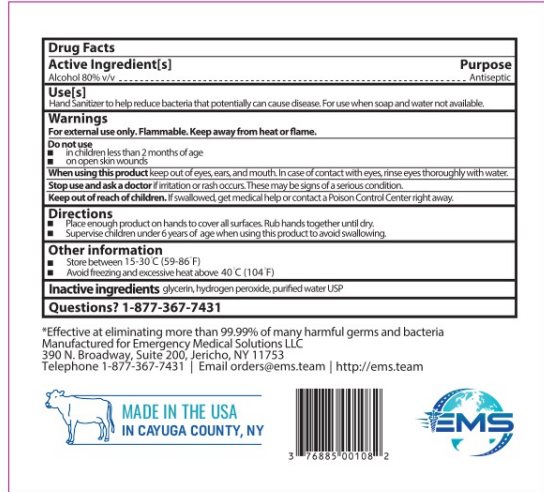


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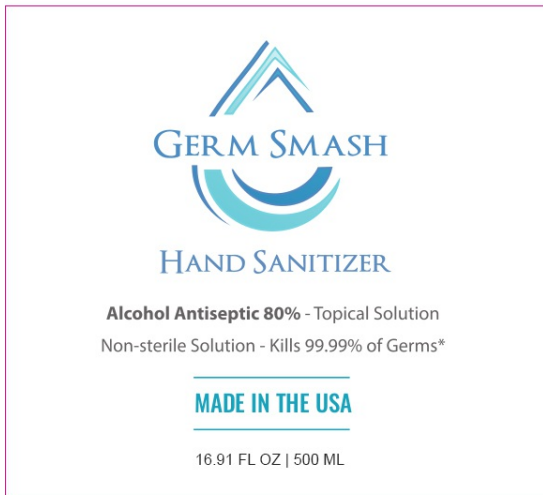


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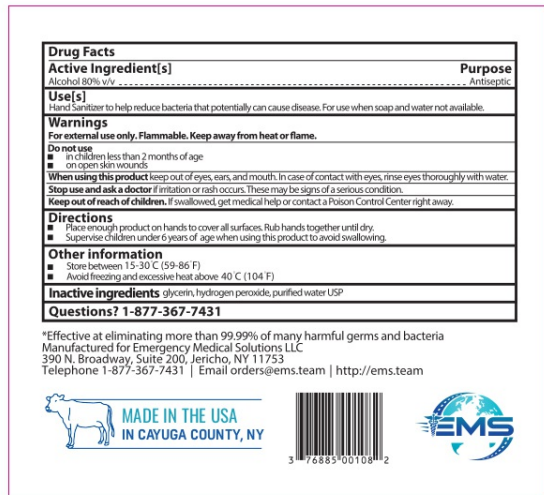


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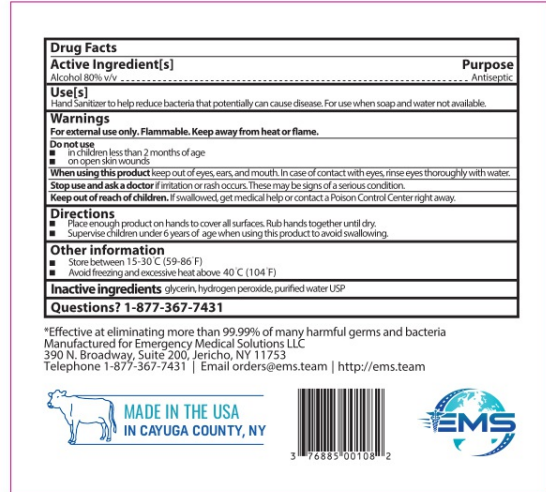


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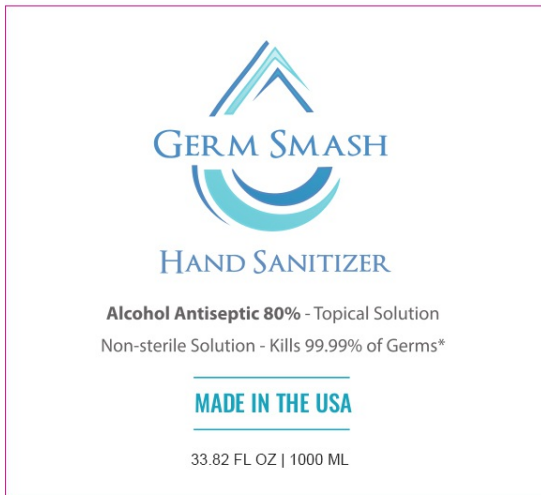


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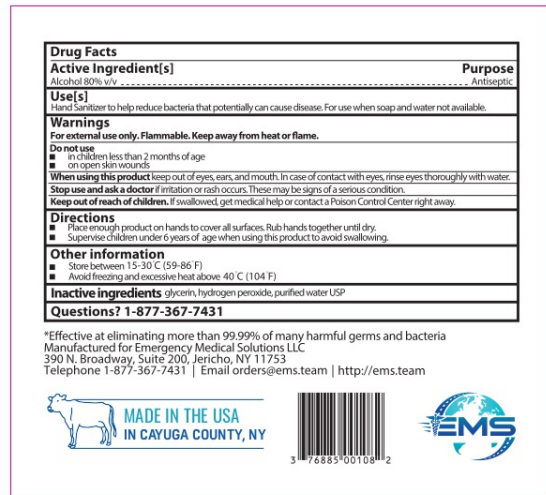


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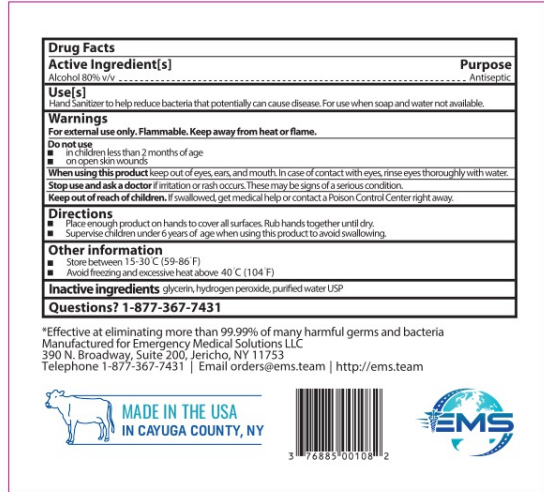


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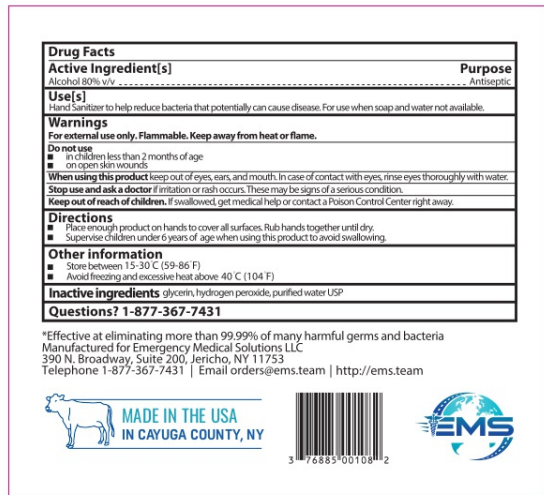


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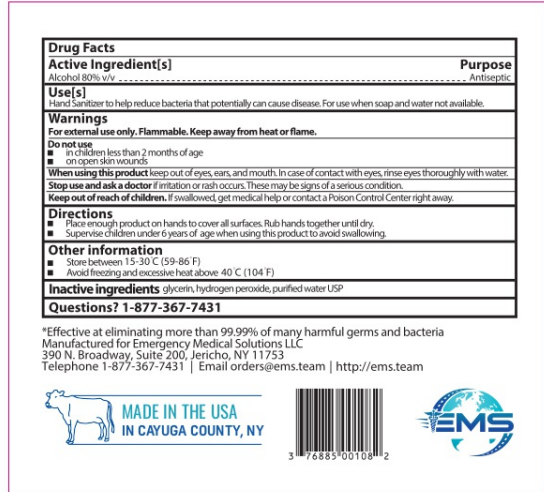


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76885-004-11

HAND SANITIZER				
alcohol liquid				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76885-004	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)		ALCOHOL	80 mL in 100 mL	
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: 059QF0K00R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76885-004-01	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

2	NDC:76885-004-02	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
3	NDC:76885-004-03	236.58 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
4	NDC:76885-004-04	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
5	NDC:76885-004-05	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
6	NDC:76885-004-06	754.125 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
7	NDC:76885-004-07	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
8	NDC:76885-004-08	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
9	NDC:76885-004-09	208198 mL in 1 DRUM; Type 0: Not a Combination Product	03/30/2020	
10	NDC:76885-004-10	1040988.24 mL in 1 CONTAINER; Type 0: Not a Combination Product	03/30/2020	
11	NDC:76885-004-11	18927.1 mL in 1 BOX; 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** - Emergency Medical Solutions LLC (117478411)

## Establishment

Name	Address	ID/FEI	Business Operations
Cayuga Ingredients, LLC		081361197	manufacture(76885-004)

## Establishment

Name	Address	ID/FEI	Business Operations
Emergency Medical Solutions LLC		117478411	label(76885-004)

## Establishment

Name	Address	ID/FEI	Business Operations
Azure Bottling, LLC.		624193384	repack(76885-004)

Revised: 6/2020

Emergency Medical Solutions LLC