HAND SANITIZER- alcohol liquid Riedel Marketing Group

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.

Germ Shield 80 Hand Sanitizer

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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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.

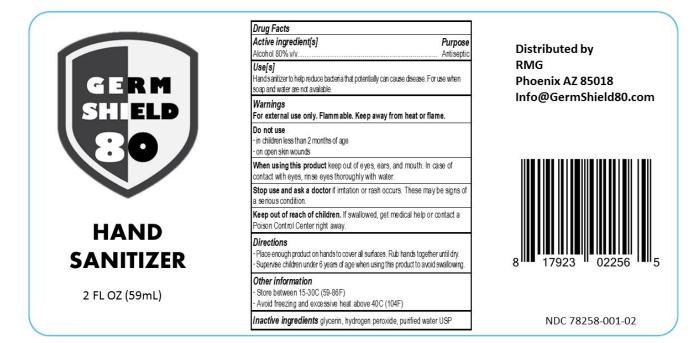
Other information

- 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



Border indicates edge of label and does not print

59 mL NDC: 78258-001-02

HAND SANITIZER			
alcohol liquid			
Product Information			
Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:78258-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety						
Ingredient Name		Basis of Strength		Strength		
ALCOHOL (UNII: 34	K9958'	V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL		80 mL in 100 mL	
Inactive Ingred	ients					
Ingredient Name				Strength		
GLYCERIN (UNII: PDC6A3C0OX)				1.45 mL in 100 mL		
HYDROGEN PEROXIDE (UNII: BBX060AN9V)				0.125 mL in 100 mL		
WATER (UNII: 059QF0KO0R)						
Packaging						
# Item Code		Package Description	Marketing Start Date		Marketing End Date	
1 NDC:78258-001- 02	59 m Produ	L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination act	05/18/2020			
	_					
Marketing Information						
Marketing Categ	ory	Application Number or Monograph Citation	Market	ing Start Date	Marketing End Date	
OTC monograph not	final	part333A	05/18/2020			

Labeler - Riedel Marketing Group (797745564)

Registrant - Flavorcraft Industries, Inc. (621632921)

Establishment

Name	Address	ID/FEI	Business Operations
Flavorcraft Industries, Inc.		621632921	manufacture(78258-001)

Revised: 6/2020

Riedel Marketing Group