HAND SANITIZER- alcohol gel Fareva Morton Grove, Inc.

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.

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)

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (78%, 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. Glycerin (1.72% v/v).
- c. Hydroxypropylcellulose (1.28% v/v).
- d. Sterile distilled water or boiled cold water.

Active Ingredient(s)

Alcohol 78% 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.

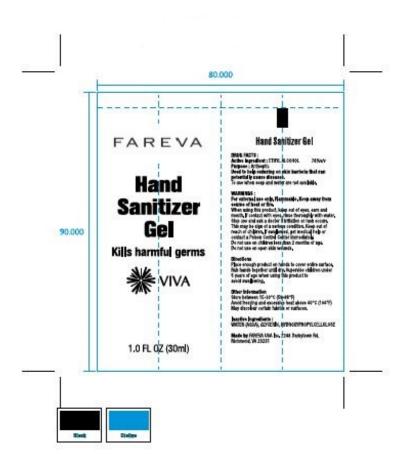
Other information

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

Inactive ingredients

Water (Aqua), Glycerin, Hydroxypropylcellulose

Package Label - Principal Display Panel



30 mL NDC: 72686-180-01

HandSanitizerproof9.jpg

HAND SANITIZER alcohol gel						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:726		NDC:72686-180		
Route of Administration	TOPICAL					
Active Ingredient/Active Moiety						
Ingredient Name			Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	78 mL in 100 mL		

Inactive Ingredients							
Ingredient Name							
GLYCERIN (UNII: PDC6A3C0OX)							
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)							
WATER (UNII: 059QF0KO0R)							
Packaging # Item Code Package Description Marketing Start Date							
0		Marketing End Date					
mL in 1 TOBE; Type 0: Not a Combination Product	04/20/2020						
Marketing Information							
Application Number or Monograph Citation		Marketing End Date					
part333A	04/20/2020						
	Ingredient Name COOX) ULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH) OR) Package Description mL in 1 TUBE; Type 0: Not a Combination Product mation Application Number or Monograph Citation	Ingredient Name 1 COOX) 1 ULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH) 1 OR) 0 Marketing Start Date Package Description mL in 1 TUBE; Type 0: Not a Combination Product 04/20/2020 Imation Marketing Start Date					

Labeler - Fareva Morton Grove, Inc. (116752326)

Establishment

Name	Address	ID/FEI	Business Operations
Fareva Morton Grove, Inc.		116752326	manufacture(72686-180), analysis(72686-180), label(72686-180), pack(72686-180), relabel(72686-180), repack(72686-180)

Revised: 4/2020

Fareva Morton Grove, Inc.