HAND SANITIZER- alcohol liquid Ida Graves 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

Drug Facts	
Active ingredient[s] Alcohol 80% v/v	Purpose
	Апизерис
Use[s] Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when	n 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 ey	es, rinse eyes thoroughly with water.
Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious of	ondition.
Keep out of reach of children. If swallowed, get medical help or contact a Poison Cont	rol 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	ş.
Other information	
 Store between 15-30C (59-86F) 	
 Avoid freezing and excessive heat above 40C (104F) 	
Inactive ingredients glycerin, hydrogen peroxide, purified water USP	

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Antiseptic, Hand Sanitizer

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Package Label - Principal Display Panel

Alcohol Antiseptic 80%

Topical Solution

Hand Sanitizer

Non-sterile Solution

3785.41 mL



Alcohol Antiseptic 80% Topical Solution Hand Sanitizer Non-sterile Solution 3785.41 mL

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

HAND SANITIZER alcohol liquid				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:74392-0001		NDC:74392-0001
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	

ALCOHOL

141.6 mL in 177 mL

Inactive Ingredients			
Ingredient Name	Strength		
GLYCEROL FORMAL (UNII: 3L7GR2604E)	2.57 mL in 177 mL		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.23 mL in 177 mL		
WATER (UNII: 059QF0KO0R)	32.6 mL in 177 mL		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:74392- 0001-1	3785.41 mL in 1 BOTTLE, PLASTIC; 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 - Ida Graves LLC (123398127)

Establishment				
Name	Address	ID/FEI	Business Operations	
Ida Graves, LLC		123398127	manufacture (74392-0001)	

Revised: 4/2020 Ida Graves LLC