HAND SANITIZER- ethyl alcohol aerosol TAGBUFF 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.

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. Isopropyl Alcohol (75%, 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)

Isopropyl Alcohol 75% 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





60ml NDC: 74927-641-60

HAND SANITIZER						
ethyl alcohol aerosol						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Sour	ce)	NDC:749	27-641	
Route of Administration	TOPICAL					
Active Ingredient/Active Moiety						
Ing	redient Name		Basis of Stre	ength	Strength	

ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL -	ISOPROPYL	75 mL
UNII:ND2M416302)	ALCOHOL	in 100 mL

Inactive Ingredients				
Ingredient Name	Strength			
GLYCERIN (UNII: PDC6 A3C0 OX)	4 mL in 100 mL			
TRICLOSAN (UNII: 4NM5039 Y5X)	3 mL in 100 mL			
WATER (UNII: 059QF0KO0R)				

Product Characteristics				
Color	Score			
Shape	Size			
Flavor	Imprint Code			
Contains				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:74927-641-60	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020		
2	NDC:74927-641-51	500 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 - TAGBUFF INC. (117472155)

Establishment				
Name	Address	ID/FEI	Business Operations	
TAGBUFF INC.		117472155	manufacture(74927-641)	

Revised: 4/2020 TAGBUFF INC.