HAND SANITIZER- alcohol solution Old Dominick Distillery

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.

Dominick 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

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Package Label - Principal Display Panel

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HAND SANITIZER

alcohol solution

Product Type	HUMAN OTC DRUG	Item Code (Sour	rce) N	DC:74049-001				
Route of Administration	TOPICAL							
Active Ingredient/Active Moiety								
	Ingredient Name	Bas	is of Strength	Strength				
ALCOHOL (UNII: 3K9958V9	Ingredient Name 00M) (ALCOHOL - UNII:3K9958V90M)	Bas ALCO	•	Strength 80 mL in 100 mL				
ALCOHOL (UNII: 3K9958V9 Inactive Ingredients	•		•					
	•		HOL					
	00M) (ALCOHOL - UNII:3K9958V90M) Ingredient Name		HOL	80 mL in 100 mL				
Inactive Ingredients	OOM) (ALCOHOL - UNII:3K9958V90M) Ingredient Name		HOL	80 mL in 100 mL trength mL				

Packaging									
#	Item Code	tem Code Package Description Marketing Start Date		Marketing End Date					
1	NDC:74049-001- 01	118.3 mL in 1 BOTTLE; Type 0: Not a Combination Product		03/30/2020					
2	NDC:74049-001- 02	118.3	B mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020					
3	NDC:74049-001- 03	177.4	4 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020					
4	NDC:74049-001- 04	1892 Pro d	.71 mL in 1 BOTTLE; Type 0: Not a Combination uct	03/30/2020					
Marketing Information									
Marketing Category		ry	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not fina		nal	part333A	03/30/2020					

Labeler - Old Dominick Distillery (876687927)

Establishment

Name	Address	ID/FEI	Business Operations
Old Dominick Distillery		876687927	manufacture(74049-001)

Revised: 3/2020

Old Dominick Distillery