

HAND SANITIZER- ethyl alcohol liquid
Mills tone Spirits Group 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.

Hand sanitizer

Alcohol 80 % v/v

Antiseptic

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

For external use only. Flammable. Keep away from heat or flame

- In children less than 2 months of age
- On open skin wounds

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.

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 40 C (104F)

Inactive ingredients glycerin, hydrogen peroxide, purified water USP

Hand Sanitizer

Hand Sanitizer Front Label

New Liberty Distillery Philadelphia PA

Alcohol

Glycerin

water

HAND SANITIZER

New Liberty Distillery
Philadelphia, PA

Ethanol 96%; Hydrogen Peroxide 3%
Glycerol 98%

ALCOHOL ANTISEPTIC SOLUTION
TOPICAL SOLUTION
NON-STERILE SOLUTION
8 ounces

hydrogen peroxide

8 ounces

HAND SANITIZER

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73887-236
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety					
		Ingredient Name	Basis of Strength	Strength	
		ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL	
Inactive Ingredients					
		Ingredient Name	Strength		
		WATER (UNII: 059QF0KO0R)			
		GLYCERIN (UNII: PDC6A3C0OX)			
		HYDROGEN PEROXIDE (UNII: BBX060AN9V)			
Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:73887-236-02	10000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/28/2020		
Marketing Information					
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final		part333A	03/28/2020		

Labeler - Millstone Spirits Group LLC (012809011)

Registrant - Millstone Spirits Group LLC (012809011)

Establishment

Name	Address	ID/FEI	Business Operations
Millstone Spirits Group LLC		012809011	manufacture(73887-236)

Revised: 3/2020

Millstone Spirits Group LLC