HAND SANITIZER- alcohol liquid New Holland Brewing Company, 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

Alconol 80% W

Use[s]

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 a sign 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 40C (104F)

Inactive Ingredients glycerin, hydrogen peroxide, denatonium benzoate, purified water USP

WARNING: KEEP OUT OF REACH OF CHILDREN

Purpose

Antiseptic, Hand Sanitizer

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Package Label - Principal Display Panel

3785 mL NDC: 74258-0008-1





ALCOHOL ANTISEPTIC 80% TOPICAL SOLUTION

> Non-Sterile Solution 1 GALLON (3.78L)

Drug Facts

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Purpose .. Antiseptic

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Other information
Sions between 15–30C (69–88F)
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Inactive Ingredients glycerin, hydrogen perceide, denatorium benzoale, purified water USP

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MANE BY NEW BOLLAND BREWING COMPANY BOLLAND, NO - BLADE IN THE USA NDC: 74258-0008



273 mL NDC: 74258-0008-2



ALCOHOL ANTISEPTIC 80% TOPICAL SOLUTION



Non-Sterile Solution 8 FL. OZ. (236mL)

Drug Facts

Active Ingredient[s] Alcohol 80% v/v.....

Purpose ..Antiseptic

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

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MARIE BY NEW HOLLAND BREWING COMPANY HOLLAND, MI - MADE IN THE USA

NDC: 74258-0008



HAND SANITIZER

alcohol liquid

Product Information

HUMAN OTC DRUG NDC:74258-0008 Product Type Item Code (Source)

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	3028 mL in 3785 mL

Inactive Ingredients Ingredient Name Strength **GLYCERIN** (UNII: PDC6A3C0OX) 64.35 mL in 3785 mL HYDRO GEN PERO XIDE (UNII: BBX060AN9V) 4.73 mL in 3785 mL

WATER (UNII: 059QF0KO0R)	681 mL in 3785 mL
DENATO NIUM BENZO ATE (UNII: 4YK5Z54AT2)	17.71 mg in 3785 mL

l	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:74258- 0008-1	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020		
	2 NDC:74258- 0008-2	237 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 - New Holland Brewing Company, LLC (179932785)

Establishment				
Name	Address	ID/FEI	Business Operations	
New Holland Brewing Company, LLC		179932785	manufacture(74258-0008)	

Revised: 4/2020 New Holland Brewing Company, LLC