

SURFACE PREP 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:

- 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.
- Glycerol (1.45% v/v).
- Hydrogen peroxide (0.125% v/v).
- 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]	Purpose
Alcohol 80% v/v	Antiseptic

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: 74258-0007



SurfacePrep

**HAND
SANITIZER**

Non-Sterile Solution

**ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION**

1 GALLON (3.78L)

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MADE BY NEW HOLLAND
BREWING COMPANY
HOLLAND, MI

NDC:74258-0007



6 14036 24601 3

237 mL NDC: 74258-0007



HAND SANITIZER

Non-Sterile Solution

**ALCOHOL ANTISEPTIC 80%
TOPICAL SOLUTION**

8 FL. OZ. (236mL)

Drug Facts

Active Ingredient(s) Alcohol 80% v/v.....	PurposeAntiseptic
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Use(s)
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Warnings
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Stop use and seek a doctor if irritation or rash occurs. There 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 8 years of age when using this product to avoid swallowing.

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KEEP OUT OF REACH OF CHILDREN

MADE BY
NEW HOLLAND BREWERS COMPANY
HOLLAND, MI • MADE IN THE USA
NDC: 74258-0007



SURFACE PREP HAND SANITIZER

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74258-0007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	3028 mL in 3785 mL

Inactive Ingredients

Ingredient Name	Strength
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GLYCERIN (UNII: PDC6A3C0OX)	64.35 mL in 3785 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	4.73 mL in 3785 mL
WATER (UNII: 059QF0K00R)	681 mL in 3785 mL
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	17.71 mg in 3785 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74258-0007-1	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:74258-0007-2	237 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 - New Holland Brewing Company, LLC (179932785)

Establishment

Name	Address	ID/FEI	Business Operations
New Holland Brewing Company, LLC		179932785	manufacture(74258-0007)

Revised: 4/2020

New Holland Brewing Company, LLC