

## **NIPPITATY DISTILLERY HAND SANITIZER- nippitaty distillery hand sanitizer liquid Traxler Littlejohn Enterprises 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.*

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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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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**

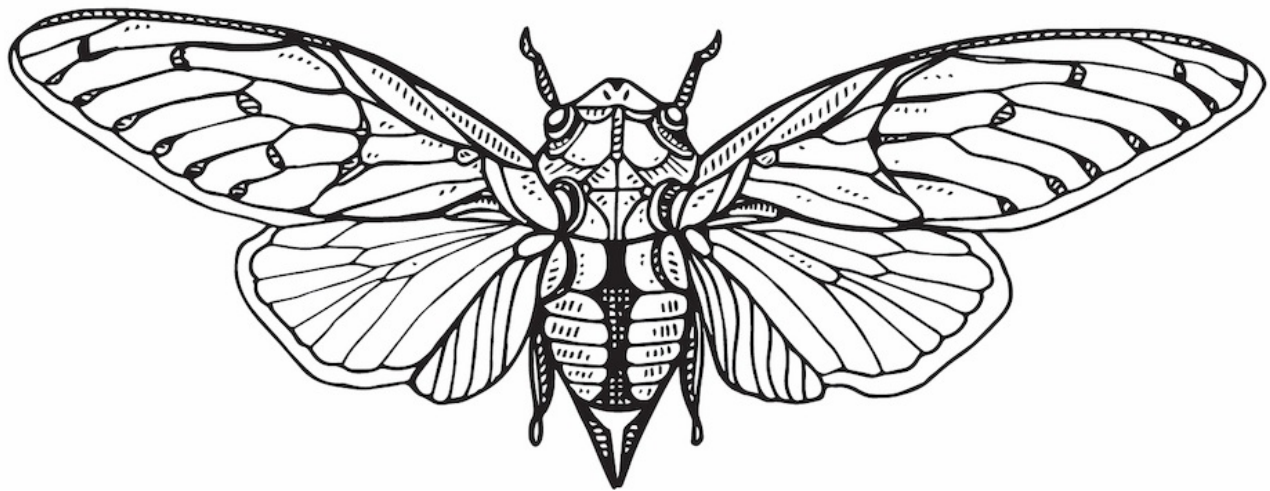
- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

**Inactive ingredients**

glycerin, hydrogen peroxide, purified water USP

**Package Label - Principal Display Panel**

NIPPITATY DISTILLERY  
— EST. 2017 —



3.66L NDC: 74343-001-02

Hand Sanitizer

Alcohol Antiseptic 80%

Topical Solution

WHO Recommended Hand Rub

Formulation #1

FOR EXTERNAL USE ONLY

NOT SAFE FOR CONSUMPTION

NON-STERILE SOLUTION

NET WT: 5GAL (17981ml)

Traxler Littlejohn Enterprises LLC (DBA Nippitaty Distillery)

4208 Pace St North Charleston SC 29405

Particularly Good and Strong Sanitizer

Made In USA

Natural Separation May Occur...Shake Well

## NIPPITATY DISTILLERY HAND SANITIZER

nippitaty distillery hand sanitizer liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:74343-001
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	15142 L in 18317 L

### Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	261 L in 18317 L
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	749 L in 18317 L
WATER (UNII: 059QF0KO0R)	2165 L in 18317 L

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74343-001-01	18317 L in 1 CONTAINER; Type 0: Not a Combination Product	03/30/2020	
2	NDC:74343-001-02	3.66 L in 1 CONTAINER; Type 0: Not a Combination Product	03/30/2020	
3	NDC:74343-001-03	1.89 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020	
4	NDC:74343-001-04	1.89 L in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/30/2020	
5	NDC:74343-001-05	0.236 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2020	
6	NDC:74343-001-06	208 L in 1 CONTAINER; Type 0: Not a Combination Product	03/30/2020	
7	NDC:74343-001-07	1022 L in 1 CONTAINER; 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** - Traxler Littlejohn Enterprises LLC (102261180)

**Establishment**

Name	Address	ID/FEI	Business Operations
Traxler Littlejohn Enterprises LLC		102261180	manufacture(74343-001)

Revised: 4/2020

Traxler Littlejohn Enterprises LLC