

## **NON ALCOHOL HAND SANITIZER- non alcohol hand sanitizer liquid**

**Uweport 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. Benzalkonium Chloride ( 0.13% w/v).
- b. Coco-Glucoside (1.5% v/v).
- c. Laurtrimonium Chloride (0.3% v/v).
- d. Cocamidopropylamine Oxide (0.3% v/v).
- e. 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)**

Benzalkonium Chloride 0.13% w/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**

Coco-Glucoside, Laurtrimonium Chloride, Cocamidopropylamine Oxide, purified water USP

**Package Label - Principal Display Panel**



ITEM #3080L4 - NON-ALCOHOL HAND SANITIZER

# Touché™ FOAMING NON-ALCOHOL HAND SANITIZER

Amano Pioneer Eclipse® Corp.  
1-800-367-3550 • 1-336-372-8080  
www.pioneereclipse.com  
1 Eclipse Road, PO Box 909  
Sparta, NC 28675 • USA  
Fax: 1-336-372-2895

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## Drug Facts

Active Ingredient .....	Purpose
Benzalkonium Chloride 0.13% .....	Antimicrobial

**Uses** ■ For handwashing to decrease bacteria on the skin. Recommended for repeat use.

### Warnings

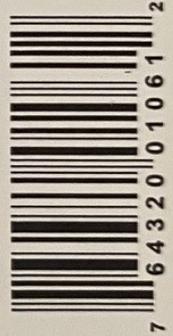
For external use only.

**Avoid contact with eyes** - In case of eye contact, flush eyes with water.

**Keep out of reach of children.** If swallowed, get immediate medical attention. Stop use and ask doctor if irritation or redness develops and persists.

**Directions** Apply foam sanitizer to hands. Rub over surfaces of both hands for 15 seconds. No rinsing required.

**Inactive ingredients** Water, Coco-Glucoside, Laurtrimonium Chloride, Cocamidopropylamine Oxide, Citric Acid, Fragrance.



## NON-ALCOHOL HAND SANITIZER

CONTENTS:  
1 Liter (33.8 fl. oz.)



1000 ml NDC: 80401-102-10

### NON ALCOHOL HAND SANITIZER

non alcohol hand sanitizer liquid

#### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80401-102
Route of Administration	TOPICAL		

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	<b>BENZALKONIUM CHLORIDE</b>	0.13 g in 100 mL
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### Inactive Ingredients

Ingredient Name	Strength
<b>COCAMIDOPROPYLAMINE OXIDE</b> (UNII: M4SL82J7HK)	0.3 mL in 100 mL
<b>LAURTRIMONIUM CHLORIDE</b> (UNII: A81MS10FIC)	0.3 mL in 100 mL
<b>COCO GLUCOSIDE</b> (UNII: ICS790225B)	1.5 mL in 100 mL
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80401-102-10	1000 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** - Uweport LLC (081252924)

**Registrant** - Uweport llc (081252924)

### Establishment

Name	Address	ID/FEI	Business Operations
Uweport		081252924	label(80401-102)

### Establishment

Name	Address	ID/FEI	Business Operations
Guangdong Bolicen Bio-Technology Co Ltd		554525110	manufacture(80401-102)

Revised: 10/2020

Uweport LLC