**ANTIBACTERIAL HAND WASH- benzalkonium chloride 0.13% liquid**
**Uline**

*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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**Antibacterial Hand Soap**
**488.003/488AD**

## Active ingredient

Benzalkonium chloride 0.13%

## Purpose

Antibacterial

## Use

for handwashing to decrease bacteria on the skin

## Warnings

For external use only: hands

## When using this product

- avoid contact with the eyes.  If contact occurs, rinse eyes with water

## Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

## Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

## Directions

- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

**Inactive ingredients**

water, lauramine oxide, cocamidopropyl betaine, lauramidopropylamine oxide, sodium chloride, myristamidopropylamine oxide, glycerin, fragrance, distearetth-75 IPDI, PEG-150 distearate, citric acid, tetrasodium EDTA, benzophenone-4, sodium benzoate, red 33, red 40, yellow 5

Distributed by: ULINE, 12575 Uline Drive

Pleasant Prairie, WI 53158

1-800-295-5510

unline.com

**principal display panel**

ULINE

HAND SOAP

ANTIBACTERIAL

S-17080

1 GAL (3.78 L)

1-800-295-5510

uline.com


**ANTIBACTERIAL HAND WASH**

benzalkonium chloride 0.13% liquid

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:69790-488 |
| **Route of Administration** | TOPICAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **BENZALKONIUM CHLORIDE** (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 1.313 mg in 1 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **water** (UNII: 059QF0KO0R) | |
| **LAURAMINE OXIDE** (UNII: 4F6FC4MI8W) | |
| **COCAMIDOPROPYL BETAINE** (UNII: 5OCF3O11KX) | |
| **LAURAMIDOPROPYLAMINE OXIDE** (UNII: I6KX160QTV) | |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) | |
| **myristamidopropylamine oxide** (UNII: 3HSF539C9T) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **DISTEARETH-75 ISOPHORONE DIISOCYANATE** (UNII: 5365FJ30SC) | |
| **PEG-150 distearate** (UNII: 6F36Q0I0AC) | |
| **CITRIC ACID MONOHYDRATE** (UNII: 2968PHW8QP) | |
| **EDETATE SODIUM** (UNII: MP1J8420LU) | |
| **SULISOBENZONE** (UNII: 1W6L629B4K) | |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **D&C RED NO. 33** (UNII: 9DBA0SBB0L) | |
| **FD&C RED NO. 40** (UNII: WZB9127XOA) | |
| **FD&C YELLOW NO. 5** (UNII: I753WB2F1M) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:69790-488-08 | 3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 05/27/2018 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333A | 05/27/2018 | |

## Labeler - Uline (039612668)

## Registrant - Vi-Jon, LLC (790752542)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Vi-Jon, LLC | | 790752542 | manufacture(69790-488) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Vi-Jon, LLC | | 088520668 | manufacture(69790-488) |