**KIMBALL MIDWEST ANTIBACTERIAL HAND CLEANER- benzalkonium chloride liquid**
**Kimball Midwest**

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***KIMBALL MIDWEST* Antibacterial Hand Cleaner**

***Drug Facts***

***Active Ingredient***

Benzalkonium Chloride 0.13% w/v

***Purpose***

Antibacterial Agent

***Uses***

To decrease bacteria on the skin.

***Warnings***

**For external use only.**

**Avoid contact with eyes.** If contact occurs, flush eyes with water.

**Stop use and ask a doctor if,** in rare instances, redness or irritation develops and persists for more than 72 hours.

**Keep out of reach of children.** If swallowed, contact a physician or poison control center.

***Directions***

Apply small amount, covering hands with product for 30 seconds. Add water, lather, rinse.

**Inactive Ingredients**

Water, Cetrimonium Chloride, Lauramine Oxide, Sorbitol, Cocamide MEA, Sodium Chloride, PEG-120 Methyl Glucose Dioleate, Sodium Lauraminodipropionate, Citric Acid, Fragrance, Disodium EDTA, Methylisothiazolinone, Methychloroisothiazolinone, Yellow 5, Red 33.

**Proven To Kill A Broad Spectrum Of Germs And Bacteria**

**Mild Yet Effective**

**Pleasant Spicy Floral Fragrance**

Carefully read warnings on side panel

**FOR INDUSTRIAL AND PROFESSIONAL USE ONLY**

Manufactured for:
**Kimball Midwest**
4800 Roberts Rd.
Columbus, Ohio 43228
800-233-1294
www.kimballmidwest.com

DESCRIPTION
The Antibacterial Hand Cleaner is a fast-acting hand cleaner with a pleasant spicy floral fragrance. Kill germs without irritating the skin and helps prevent drying, cracking and chapping.

DIRECTIONS
Apply Antibacterial Hand Cleaner to wet hands. Work thoroughly into hands, knuckles and under fingernails. Rinse with water.

**Made in the U.S.A.**

**Packaging**


## KIMBALL MIDWEST ANTIBACTERIAL HAND CLEANER

benzalkonium chloride liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:66608-216 |
| **Route of Administration** | TOPICAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **BENZALKONIUM CHLORIDE** (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y) | BENZALKONIUM CHLORIDE | 1.3 mg in 1 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | |
| **CETRIMONIUM CHLORIDE** (UNII: UC9PE95IBP) | |
| **LAURAMINE OXIDE** (UNII: 4F6FC4MI8W) | |
| **SORBITOL** (UNII: 506T60A25R) | |
| **COCO MONOETHANOLAMIDE** (UNII: C80684146D) | |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) | |
| **PEG-120 METHYL GLUCOSE DIOLEATE** (UNII: YM0K64F20V) | |
| **SODIUM LAURIMINODIPROPIONATE** (UNII: 7G447D0DH9) | |
| **CITRIC ACID MONOHYDRATE** (UNII: 2968PHW8QP) | |
| **EDETATE DISODIUM** (UNII: 7FLD91C86K) | |
| **METHYLISOTHIAZOLINONE** (UNII: 229D0E1QFA) | |
| **METHYLCHLOROISOTHIAZOLINONE** (UNII: DEL7T5QRPN) | |
| **FD&C YELLOW NO. 5** (UNII: I753WB2F1M) | |
| **D&C RED NO. 33** (UNII: 9DBA0SBB0L) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:66608-216-01 | 3785 mL in 1 BOTTLE; Type 0: Not a Combination Product | 11/13/2024 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | 505G(a)(3) | 11/13/2024 | |

## Labeler - Kimball Midwest (017906231)

Revised: 11/2024 Kimball Midwest