**2PK HAND SANITIZER CLASSIC- alcohol gel**
**UNIVERSAL DISTRIBUTION CENTER LLC**

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**Active ingredient**

Ethyl Alcohol 62%

**Purpose**

Antimicrobial

**Uses**

Hand Sanitizer to help reduce bacteria on skin.

**WARNINGS**

**Flammable. Keep away from fire or flame.**

**For external use only.**

**When using this product,**

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

**Stop use and ask a doctor if**

irritation or rash appears and lasts.

**Keep out of reach of children.**

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

**Directions:**

- Place enough product in your palm to thoroughly spread on both hands and rub into the skin until dry.
- Children under 6 years of age should be supervised when using this product.

**Other Information:**

- Store below 106℉ (41℃)
- May Discolor certain fabrics or surfaces.

**Inactive ingredients**

Water (Aqua), Aloe Barbadensis Leaf Juice, Carbomer, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate (Vitamin E), Aminomethyl Propanol.





# 2PK HAND SANITIZER CLASSIC

alcohol gel

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:52000-132 |
| **Route of Administration** | TOPICAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **ALCOHOL** (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 62 mL in 100 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | |
| **CARBOMER 934** (UNII: Z135WT9208) | |
| **.ALPHA.-TOCOPHEROL ACETATE** (UNII: 9E8X80D2L0) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **ALOE VERA LEAF** (UNII: ZY81Z83H0X) | |
| **AMINOMETHYL PROPANEDIOL** (UNII: CZ7BU4QZJZ) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:52000-132-01 | 60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 01/01/2018 | |
| **2** | NDC:52000-132-12 | 120 mL in 1 BLISTER PACK; Type 0: Not a Combination Product | 01/01/2018 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M003 | 01/01/2018 | |

## Labeler - UNIVERSAL DISTRIBUTION CENTER LLC (019180459)

## Registrant - UNIVERSAL DISTRIBUTION CENTER LLC (019180459)

Revised: 1/2023 UNIVERSAL DISTRIBUTION CENTER LLC