# HAND SANITIZER 1.850Z- 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 the skin.

### **Warnings**

For external use only.

**Flammable.** Keep away from heat and flame.

### When using this product

avoid contact with face, eyes, and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

## Stop use and ask a doctor if

irritation or redness develops.

## Keep out of reach of children.

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

#### **Directions**

- wet hands thoroughly with product and rub into skin until dry.
- Children under 6 years of age should be supervised by an adult when using.

## **Inactive ingredients**

Water (Aqua), Aloe Barbadensis Leaf Juice, Carbomer, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate (Vitamin E), Aminomethyl Propanol, (FD&C Red No. 40, FD&C Yellow No. 5, D&C Red No. 33, FD&C Blue No.1)





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ITEM#56024



Made in China

Distributed by: Universal Distribution Center 96 Distribution Boulevard Edison, NJ 08817



1.85 El 07 / 54 7 ml

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ITEM#56024



### **HAND SANITIZER 1.850Z**

alcohol gel

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:52000-131

**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

FD&C RED NO. 40 (UNII: WZB9127XOA)

WATER (UNII: 059QF0KO0R)

**CARBOMER 934** (UNII: Z135WT9208)

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

**GLYCERIN** (UNII: PDC6A3C0OX)

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
ALOE VERA LEAF (UNII: ZY81Z83H0X)
D&C RED NO. 33 (UNII: 9DBA0SBB0L)
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)

FD&C BLUE NO. 1 (UNII: H3R47K3TBD)

AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)

#### **Packaging**

ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:52000- 131-01	54.7 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2018	

## **Marketing Information**

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	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