2PK HAND SANITIZER ALOE VERA- alcohol gel UNIVERSAL DISTRIBUTION CENTER LLC

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°F (41°C)
- 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, FD&C Blue No.1, FD&C Yellow No. 5











2PK HAND SANITIZER ALOE VERA

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52000-133
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)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:52000- 133-01	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2018		
2	NDC:52000- 133-12	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