

**KISS MY FACE HAND SANITIZER WITH ALOE BACTERIAL DEFENSE FRAGRANCE
FREE- benzalkonium chloride liquid
Windmill Health Products, LLC**

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.

**Kiss My Face® Hand Sanitizer with Aloe Bacterial Defense Fragrance Free
Drug Facts**

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- For antiseptic cleansing to decrease bacteria on the skin. Recommended for repeated use.

Warnings

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Ask a doctor

Stop use and consult a doctor if irritation or redness develops. If condition persists for more than 72 hours, discontinue use.

Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Do not use on children less than 2 months of age or on open skin wounds

Directions

Apply a small amount of gel into palm of hand. Rub thoroughly over all surfaces of both hands for 15 seconds.

Other information

Store tightly closed, in a cool dry place.

Inactive ingredients

Water, Aloe Barbadensis Leaf Extract, Sodium Hyaluronate, Cocos Nucifera (Coconut) Oil, Glycerin, Polyacrylamide, Tocopheryl Acetate, Propylene Glycol, Polyquaternium-7, Polysorbate 20, C13-14 Isoparaffin, Laureth-7, Phenoxyethanol, Aminomethyl Propanol

Questions or comments

1.800.822.4320

Made in the USA.

PRINCIPAL DISPLAY PANEL - 502 ML Bottle Label

KISS
MY
FACE®

HAND
SANITIZER
WITH ALOE

BACTERIAL DEFENSE

FRAGRANCE FREE

MOISTURIZING FORMULA
ANTIMICROBIAL ANTISEPTIC
ALCOHOL FREE | PARABEN FREE
17 FL OZ (502 ML)

310L0835001



KISS MY FACE HAND SANITIZER WITH ALOE BACTERIAL DEFENSE FRAGRANCE FREE

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74154-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	130 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

ALOE VERA LEAF (UNII: ZY81Z83H0X)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
COCONUT OIL (UNII: Q9L0O73W7L)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
LAURETH-7 (UNII: Z95S6G8201)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74154-006-17	502 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333A	05/08/2020	

Labeler - Windmill Health Products, LLC (831136267)

Establishment

Name	Address	ID/FEI	Business Operations
Cemi International		015038336	MANUFACTURE(74154-006)

Revised: 2/2022

Windmill Health Products, LLC