DEWZI ALCOHOL FREE HAND SANITIZER FRAGRANCE FREE- benzalkonium chloride liquid
LAB-CLEAN, INC

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.

Active ingredient

Benzalkonium Chloride 0.1%

Purpose

Antimicrobial

Uses

For hand sanitizing 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.

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

Keep out of reach of children.

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

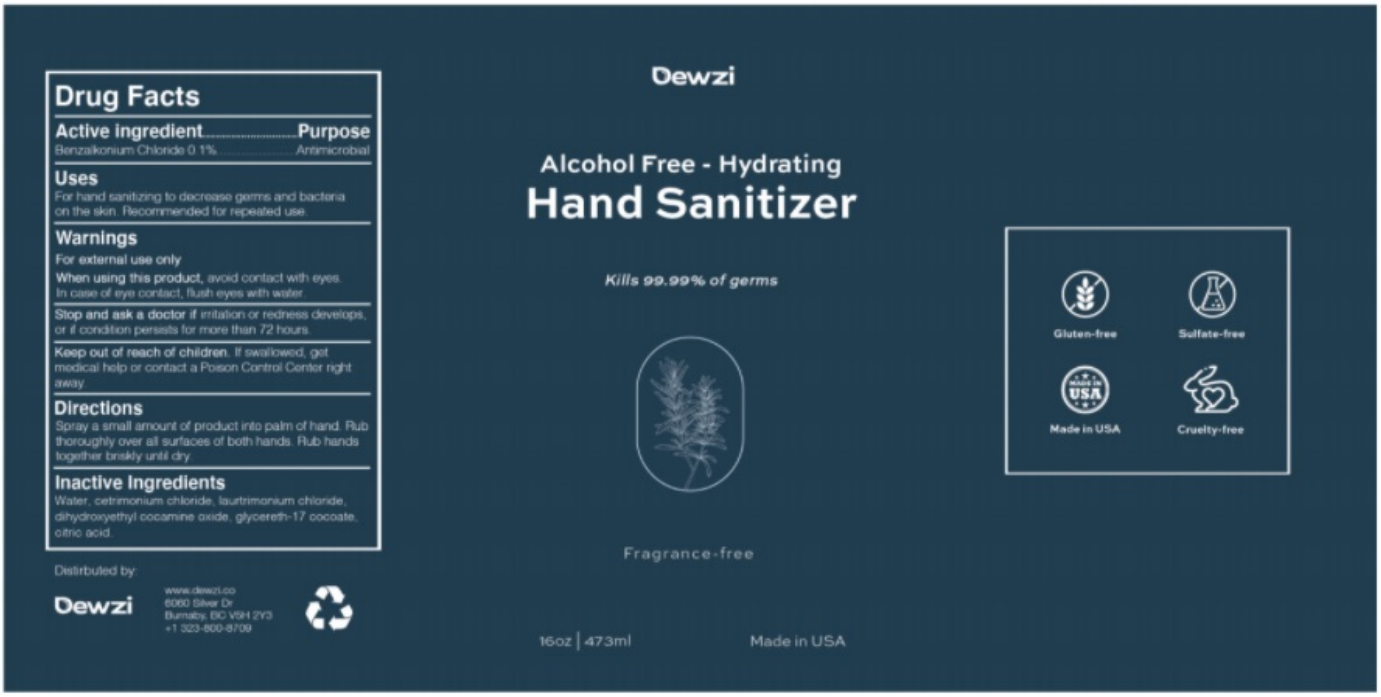
Directions

- Spray a small amount of product into the palm of hand. Rub thoroughly over all surfaces of both hands.
- Rub hands together briskly until dry.

Inactive ingredients

Water, Cetrimonium Chloride, Laurtrimonium Chloride, Dihydroxyethyl Cocamine Oxide, Glycereth-17 Cocoate, Citric Acid

Product label



DEWZI ALCOHOL FREE HAND SANITIZER FRAGRANCE FREE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73126-031
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	
GLYCERETH-17 COCOATE (UNII: 3057VPT0KC)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73126-031-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/10/2021	

2	NDC:73126-031-02	88 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/10/2021	
3	NDC:73126-031-03	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/10/2021	
4	NDC:73126-031-04	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/10/2021	
5	NDC:73126-031-05	474 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/10/2021	
6	NDC:73126-031-06	710 mL in 1 PACKAGE; Type 0: Not a Combination Product	02/10/2021	
7	NDC:73126-031-07	946 mL in 1 PACKAGE; Type 0: Not a Combination Product	02/10/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/10/2021	04/20/2024

Labeler - LAB-CLEAN, INC (199822219)

Establishment

Name	Address	ID/FEI	Business Operations
LAB-CLEAN, INC		199822219	manufacture(73126-031)

Revised: 4/2022 LAB-CLEAN, INC