

**ANTI-BACTERIAL HAND SANITIZER- benzalkonium chloride liquid
FGD, 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.

Anti-Bacterial Hand Sanitizer

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Directions

- Pump a small amount of foam into palm of hand
- Rub thoroughly over all surfaces of both hands for 15 seconds
- Rinse with potable water

Keep out of reach of children

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

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.

Uses

For hand washing to decrease bacteria on the skin

Recommended for repeated use

Inactive ingredients

Water, coco-glucoside, laurtrimonium chloride, cocamidopropylamine oxide, citric acid

Anti-bacterial Hand Sanitizer



ANTI-BACTERIAL HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73787-107
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
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CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0K00R)	
COCO GLUCOSIDE (UNII: ICS790225B)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
COCAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73787-107-01	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/11/2020	

Labeler - FGD, LLC (111927555)

Establishment

Name	Address	ID/FEI	Business Operations
Goodwin Co.		806987483	manufacture(73787-107)

Revised: 1/2021

FGD, LLC