

**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.*

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**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

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:73787-106
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

Ingredient Name	Strength
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>COCO GLUCOSIDE</b> (UNII: ICS790225B)	
<b>LAURTRIMONIUM CHLORIDE</b> (UNII: A81MSI0FIC)	
<b>COCAMIDOPROPYLAMINE OXIDE</b> (UNII: M4SL82J7HK)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73787-106-01	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(73787-106)

Revised: 1/2021

FGD, LLC