

**CONSULT HEALTH PERSISTENCE HAND SANITIZER- benzalkonium chloride liquid
Geo Management Corporation**

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.

Consult Health Persistence Hand Sanitizer

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial and First aid antiseptic

Uses:

- For hand sanitizing to decrease bacteria on the skin. Recommended for repeated use.
- For wound sanitizing to help prevent bacterial contamination in cuts, scrapes, burns, lacerations and skin infections.

Warnings:

For External Use Only.

When using this product

avoid contact with eyes. In case of eye contact, flush eyes with water.

Discontinue use if

irritation or redness develops. If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children.

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

Directions:

- Spray onto hands as needed. Rub briskly together until dry. No rinsing required.
- Spray onto wounds 3 times a day after cleaning. Allow foam to dissipate. Wipe excess with sterile gauze. May be bandaged once dry.

Inactive Ingredients

ionized water, carbamide

□Package Labeling:

CONSULT HEALTH



PERSISTENCE™
ANTI-BACTERIAL
ANTI-MICROBIAL
HAND SANITIZER

EXTENDED PROTECTION*

ELIMINATES
99.9% OF GERMS

3.0 fl oz (89 mL)

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*Studies show Benzalkonium Chloride protects against infection for more than three hours.
Distributed by: Geo Management, Las Vegas, NV 89128
Customer Service: 866-272-4425 • www.consulthealth.com

3110.0000901



CONSULT HEALTH PERSISTENCE HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70803-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
UREA (UNII: 8W8T17847W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70803-003-03	89 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/01/2020	

