BENZA CLEAN HAND SANITIZER- benzalkonium chloride liquid
BIOSYN 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 0.13%

Purpose

Antimicrobial and First Aid Antiseptic

Uses

For hand sanitizing to decrease bacteria on the skin. As a wound antiseptic to help prevent bacterial contamination in cuts, burns, scrapes, lacerations and skin infections.

Warnings

For external use only.

When using this product avoid contact with eyes. Incase of eye contact, flush eyes with water.
Discontinue useif irritation or redness develops. If condition persists for more than 72 hours, consult a doctor.

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

Directions

- Pump onto hands as needed. Rub briskly until dry.
- Apply to wounds 3 times per day after cleaning. Allow to dry. May be bandaged once dry.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Water, Sodium Bicarbonate, Colalipid C, Hydroxyethyl Cellulose, Green Tea Leaf Extract.

Package Label - Principal Display Panel

BENZA CLEAN HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79832-001
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)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:79832-001-01	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/18/2020	
2	NDC:79832-001-02	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/18/2020	

Labeler - BIOSYN INC (079797906)

Revised: 9/2020 BIOSYN INC