

**ANTIBACTERIAL- benzalkonium chloride liquid**  
**Sante Manufacturing 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.*

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Benzalkonium Chloride - 0.13%

Purpose - Antibacterial

Uses for handwashing or decrease bacteria to the skin

Warnings For external use only

Stop use and ask a doctor if irritation or redness develops

When using this product

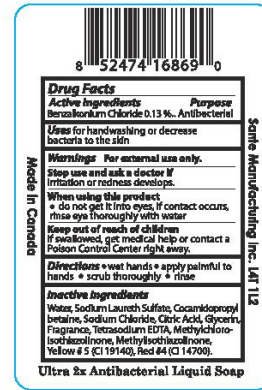
- do not get it into eyes. If contact occurs, rinse eye thoroughly with water

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

Directions

- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse

Water, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Chloride, Citric Acid, Glycerin, Fragrance, Tetrasodium EDTA, Methylchloroisoithiazolinone, Methylisoithiazolinone, Yellow# 5 (CI 19140), Red# 4 (CI 14700)



## ANTIBACTERIAL

benzalkonium chloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:71020-004
<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	1.3 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM LAURETH SULFATE</b> (UNII: BPV390UAP0)	
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)	

<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>DITETRACYCLINE TETRASODIUM EDETATE</b> (UNII: WX0A0IT7K5)	
<b>METHYLCHLOROISOTHIAZOLINONE</b> (UNII: DEL7T5QRPN)	
<b>METHYLISOTHIAZOLINONE</b> (UNII: 229D0E1QFA)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>FD&amp;C RED NO. 4</b> (UNII: X3W0AM1JLX)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71020-004-64	1900 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/08/2016	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/08/2016	

**Labeler** - Sante Manufacturing Inc (242048747)

**Registrant** - Sante Manufacturing Inc (242048747)

### Establishment

Name	Address	ID/FEI	Business Operations
Sante Manufacturing Inc		204348627	manufacture(71020-004)

### Establishment

Name	Address	ID/FEI	Business Operations
Brands International Corporation		243748238	manufacture(71020-004)

Revised: 1/2022

Sante Manufacturing Inc