

**ANTIBACTERIAL 2X ULTRA- 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.*

**Spa Soap ANTIBACTERIAL 2X- benzalkonium chloride liquid**

Benzalkonium Chloride - 0.13%

Purpose: Antibacterial

Direction

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

For external use only

Stop use and ask a doctor if irritation or redness develops

When using this product

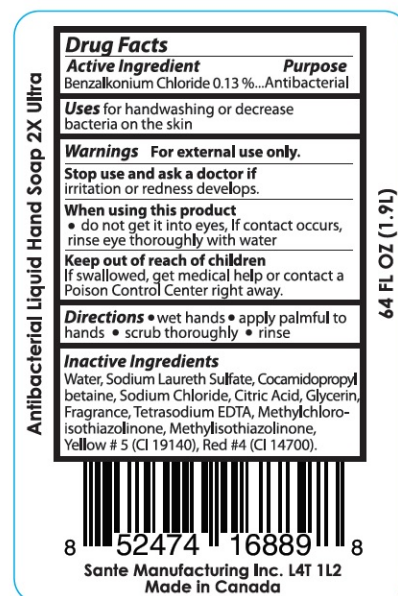
- do not get it into eyes. If contact occurs, rinse eye thoroughly with water

Keep out of reach of children

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

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

Uses for handwashing or decrease bacteria to the skin



# ANTIBACTERIAL 2X ULTRA

benzalkonium chloride liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:71020-017
<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)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71020-017-19	1900 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/08/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/08/2020	

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

Revised: 9/2020

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