

# ANTIBACTERIAL 2X- benzalkonium chloride liquid

## Sante Manufacturing Inc

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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, Methylchloro-isothiazolinone, Methylisothiazolinone, Yellow# 5 (CI 19140), Red# 4 (CI 14700)

Uses for handwashing or decrease bacteria to the skin





**SpaSoap 500 mL (16.9 oz liq.)  
Antibacterial Liquid Soap 2X  
Savon Liquide Antibactérien 2X**

**Directions:** Wet hands and apply soap. Wash vigorously and rinse. If product comes in contact with eyes, rinse with water.

**Mode d'emploi :** Mouiller les mains et appliquez du savon. Laver vigoureusement et rincer. Si le produit entre en contact avec les yeux, rincer à l'eau.

**Active Ingredient/Ingrédient tensioactif :**  
0.13% Benzalkonium Chloride

**Ingredient/Ingrédient :** Aqua, Sodium Laureth Sulfate, Sodium Chloride, Cocamidopropyl Betaine, Tetrasodium EDTA, Fragrance, Methylchloroisothiazolinone, Methylisothiazolinone, Glycerine, FD&C Red #4, FD&C Yellow #5.

Sante Manufacturing Inc.  
L4T 1L2  
Fabriqué au Canada

## ANTIBACTERIAL 2X

benzalkonium chloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:71020-016
<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

<b>1</b>	NDC:71020-016-16	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/30/2018	
<b>2</b>	NDC:71020-016-20	600 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/30/2018	

  

<b>Marketing Information</b>			
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	505G(a)(3)	01/30/2018	

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

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

<b>Establishment</b>			
<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Sante Manufacturing Inc		204348627	manufacture(71020-016)

Revised: 1/2025

Sante Manufacturing Inc