

ANTIBACTERIAL AQUA CLEAR 2X ULTRA- benzalkonium chloride liquid

Sante Manufacturing Inc

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

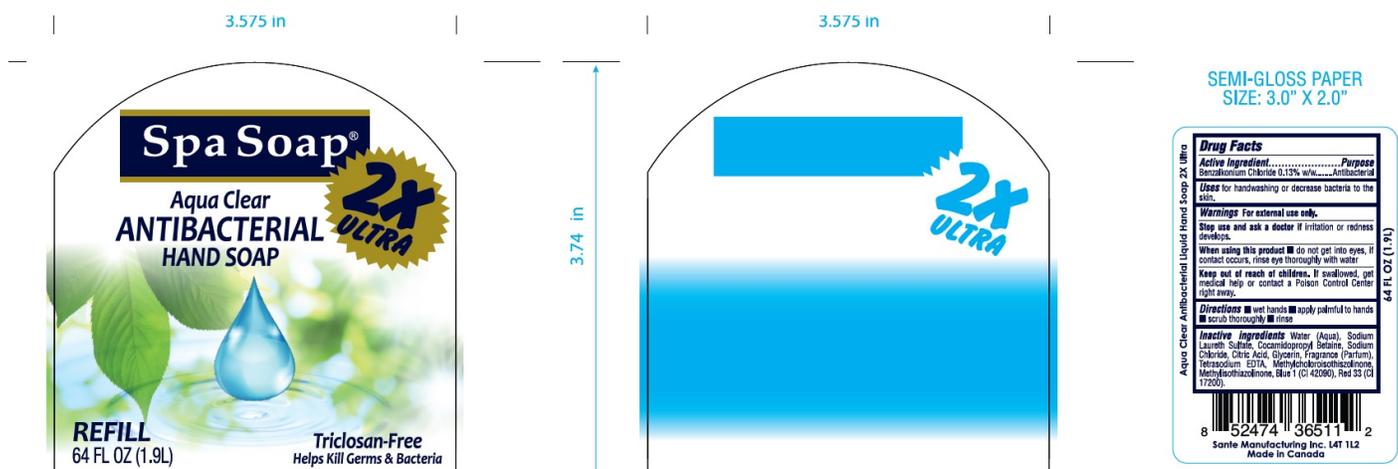
- do not get it into eyes. If contact occurs, rinse eye thoroughly with water

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, Methylchloroisothiazolinone, Methylisothiazolinone, Blue 1, Red# 4 (CI 14700)

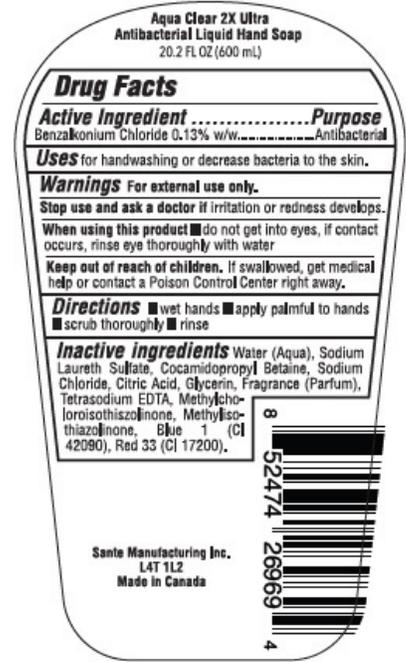
Uses for handwashing or decrease bacteria to the skin



CLEAR FILM
SIZE: 4.15" X 2.4561"



LABEL SIZE: 4.15" X 2.4561"
WHITE BOPP
BACK LABEL



ANTIBACTERIAL AQUA CLEAR 2X ULTRA

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71020-030
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
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCERIN (UNII: PDC6A3C0OX)	
DITETRACYCLINE TETRASODIUM EDETATE (UNII: WX0A0IT7K5)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	

METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)

FD&C BLUE NO. 1 (UNII: H3R47K3TBD)

FD&C RED NO. 4 (UNII: X3W0AM1JLX)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71020-030-64	1900 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/08/2016	
2	NDC:71020-030-30	600 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 Drug	505G(a)(3)	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(71020-030)

Revised: 1/2025

Sante Manufacturing Inc