SANISMART 2- benzalkonium chloride liquid
Registered Chemicals Corporation

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

Uses •For hand washing to decrease bacteria on the skin

•Recommended for repeated use

Warnings

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes thoroughly with water.

Stop use and ask a doctor if irritation or redness develop, or if condition persists for more than 72 hours.

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

Directions •Pump a small amount of foam into palm of hand.

•Rub thoroughly over all surfaces of both hands for 15 seconds. •Rinse with potable water.

Inactive ingredients Water (Aqua), coco-glucoside, laurtrimonium chloride, coamidopropylamine oxide, citric acid

Questions or comments? 781-803-7800 •info@registeredchemicals.com

SaniSmart

Antibacterial
FOAMING
HAND SO
Fragrance Free

Nonfood Compounds
Program Listed (E2)

1 US GALLON (3.785 L)

SaniSmart 2

Drug Facts

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Please Recycle

Made in the U.S.A.

Manufactured for Registered Chemicals Corporation
Weymouth, MA 02189

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SANISMART 2

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72130-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	.13 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
COCO GLUCOSIDE (UNII: ICS790225B)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
COCAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72130-002-02	1250 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/01/2018	
2	NDC:72130-002-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/01/2018	

Labeler - Registered Chemicals Corporation (046652831)

Revised: 3/2018 Registered Chemicals Corporation