DERMATIZE- benzethonium chloride liquid
Anderson Chemical Company

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

Benzethonium chloride, 0.2% w/v

Purpose

Antiseptic

Use

For handwashing to decrease bacteria on skin.

Warning

For external use only.

When using this product

- discontinue use if irritatio and redness develop.
- do not use in or near eyes.

Stop use and ask a doctor if

Stop use and ask a doctor if irritation or redness develops, or if conditions persist for more than 72 hours.

Keep out of reach of children.

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- In case of accidental ingestion, see medical attention or contact a poision control cernter immediately.

Directions

- Hands need not be washed prior to using.
- For one step hand wash/antiseptic skin cleaning.
- Place 5 milliliters in palm of hand, add water, work up a lather and scrub for 30 seconds.
- Rinse hands thoroughly with potable water after washing.

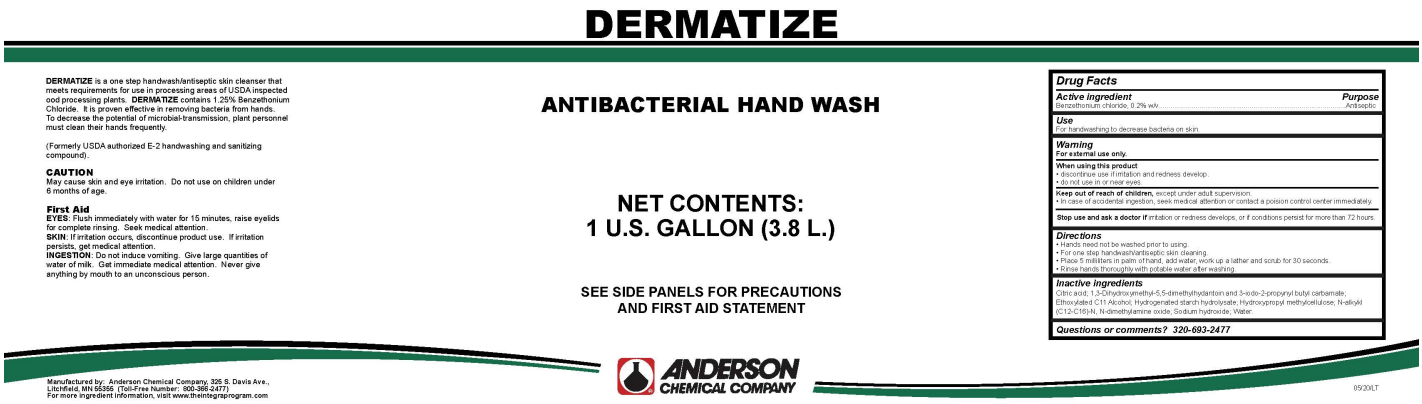
Inactive ingredients

Citric acid; 1,3-Dihydroxymethyl-5,5-dimethylhydantoin and 3-iodo-2-propynyl butyl carbamate; Ethoxylated C11 Alcohol; Hydrogenated starch hydrolysate; Hydroxypropyl methylcellulose; N-alkykl (C12-C16)-N, N-dimethylamine oxide; Sodium hydroxide; Water.

Questions or comments?

320-693-2477

Principal Display Panel



DERMATIZE

benzethonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63131-0004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	2 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
HYDROXYMETHYL CELLULOSE (UNII: 273FM27VK1)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
HYDROGENATED STARCH HYDROLYSATE (UNII: 27F77DSJ5V)	
UNDECETH-7 (UNII: R6B5PCO2JN)	
WATER (UNII: 059QF0KO0R)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63131-0004-1	15140 in 1 CASE	10/02/2020	
1		3785 mL in 1 JUG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/02/2020	

Labeler - Anderson Chemical Company (006179220)

Establishment

Name	Address	ID/FEI	Business Operations
Anderson Chemical Company		006179220	manufacture(63131-0004)

Revised: 5/2020 Anderson Chemical Company