

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 irritation 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.

Keep out of reach of children, except under adult supervision.

- In case of accidental ingestion, see medical attention or contact a poison control center 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-alkyl (C12-C16)-N, N-dimethylamine oxide; Sodium hydroxide; Water.

Questions or comments?

320-693-2477

Principal Display Panel

DERMATIZE is a one step handwash/antiseptic skin cleanser that meets requirements for use in processing areas of USDA inspected and processing plants. DERMATIZE contains 1.25% Benzethonium Chloride. It is proven effective in removing bacteria from hands. To decrease the potential of microbial transmission, plant personnel must clean their hands frequently.

(Formerly USDA authorized E-2 handwashing and sanitizing compound).

CAUTION
May cause skin and eye irritation. Do not use on children under 6 months of age.

First Aid
EYES: Flush immediately with water for 15 minutes, raise eyelids for complete rinsing. Seek medical attention.
SKIN: If irritation occurs, discontinue product use. If irritation persists, get medical attention.
INGESTION: Do not induce vomiting. Give large quantities of water or milk. Get immediate medical attention. Never give anything by mouth to an unconscious person.

ANTIBACTERIAL HAND WASH


NET CONTENTS:
1 U.S. GALLON (3.8 L.)

SEE SIDE PANELS FOR PRECAUTIONS
AND FIRST AID STATEMENT

Drug Facts

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Use For handwashing to decrease bacteria on skin.	
Warnings For external use only.	
When using this product • Discontinue use if irritation and redness develops. • Do not use in or near eyes.	
Keep out of reach of children , except under adult supervision.	
Stop use and ask a doctor if irritation or redness develops, or if conditions persist for more than 72 hours.	
Directions • Hands need not be washed prior to using. • For one step handwash/antiseptic skin cleaning. • Shake 2 milliliters in palm of hand, add water, mix up a lather and scrub for 20 seconds. • Rinse hands thoroughly with potable water after washing.	
Inactive ingredients Citric acid, 1,2-Cyclohexanediyl 5,5-dimethylphosphonate and 5-methyl-2-methyl-2-oxo-1,3-dioxane-5-carboxylate, Ethanol, Hydrogenated starch hydrolysate, Hydroxyethyl methylcellulose, Hydrolyzed C12-C18 fatty acids, Sodium hydroxide, Water.	
Questions or comments? 320-693-2477	

Manufactured by: Anderson Chemical Company, 305 S. Davis Ave.,
Litchfield, MN 55646 (Toll-free Number: 800-596-2477)
For more ingredient information, visit www.theingredientprogram.com



0929LT

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