

ASSURE- 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

Antimicrobial

Use

- For hand washing to decrease bacterial on skin.
- Recommended for repeated use.

Warning

For external use only.

When using this product

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

Stop use and ask a doctor if

Stop use and ask a doctor if irritation or redness develops, or if conditions persist.

Keep out of reach of children

Keep out of reach of children, except under adult supervision. If swallowed, get medical help or contact a Poison Control Center.

Directions

- Hands need not be washed prior to using.
- For one step handwash/antibacterial skin cleaning.
- Place product in palm of hand, add water, work up a lather.
- Rinse hands thoroughly with potable water after washing.

Inactive ingredients

C9-11 Pareth-6; Caprylyl/Capryl Oligpoglucoside; Citric acid; DMDM Hydantoin; Dye; Fragrance; Glycerine; Poly(Laurylglucoside)-7; Water.

Questions or comments?

320-693-2477

Principal Display Panel

ASSURE

Based on currently available data, this product does not meet the regulatory definition of a hazardous substance according to CSHS (HACM 2012).

Caution: May cause eye irritation.

First Aid

EYES: Flush immediately with water for 15 minutes, raise eyelids for complete rinsing. If irritation persists, call a physician.

SKIN: If irritation occurs, discontinue use. If irritation persists, call a physician.

INGESTION: Drink 1-2 glasses of water. Call a physician.

PHOSPHATE FREE

ANTIMICROBIAL FOAMING HAND SOAP

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

SEE SIDE PANELS FOR PRECAUTIONS
AND FIRST AID STATEMENT

Drug Facts	
Active Ingredient	Purpose
Benzethonium chloride, 0.2% w/v	Antimicrobial
Use	
• For hand washing to decrease bacteria on 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 with water.	
Stop use and ask a doctor if irritation or redness develops, or if conditions persist.	
Keep out of reach of children, except under adult supervision. If swallowed, get medical help or contact a poison control center right away.	
Directions	
• Hands need not be washed prior to using.	
• For one (one) use only (do not rub into skin).	
• Place product in palm of hand, add water, work up a lather.	
• Rinse hands thoroughly with potable water after washing.	
Inactive Ingredients	
C9-11 Pareth-6, Capryl/Capryl Oligoglycoside, Citric acid, DMDM Hydantoin, dye, yellow 05 liquid, dye, blue 0409 liquid, Fragrance, Glycerin, Polyoxyethylene-10-15 Sulfate, Water.	
Questions or comments? 320-693-2477	

SURFLEX

Manufactured by: Anderson Chemical Company, 326 S. Davis Ave., Litchfield, MN 55358 (Toll-Free Number: 800-366-2477)
For more ingredient information, visit www.theinfoingredient.com

060120AT

ASSURE

benzethonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63131-1132
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
DMDM HYDANTOIN (UNII: BYR0546TOW)	
C9-11 PARETH-6 (UNII: KCE0V8JT7W)	
CAPRYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLY(LAURYLGLUCOSIDE)-7 (UNII: VB00RDE21R)	
WATER (UNII: 059QF0KO0R)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63131-1132-1	7570 in 1 CASE	04/17/2007	
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	04/17/2007	

Labeler - Anderson Chemical Company (006179220)

Establishment

Name	Address	ID/FEI	Business Operations
Anderson Chemical Company		006179220	manufacture(63131-1132)

Revised: 5/2020

Anderson Chemical Company