ECOLAB - benzalkonium chloride solution
Kay 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.

Drug Facts

Active ingredient

Benzalkonium chloride 0.1%

Purpose

Antiseptic handwash

Uses

- For handwashing to decrease bacteria on the skin

Warnings

For external use only

Do not use

- In eyes

When using this product

- If in eyes, rinse promptly and thoroughly with water
- Discontinue use if irritation and redness develop

Stop use and ask a doctor if

- Skin irritation and redness occurs 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

- Wash hands to remove soil
- Dispense palmful
- Spread to cover hands, rub in well
- Air dry, do not rinse or towel dry

Other Information

- for additional information, see Material Safety Data Sheet (MSDS)

- For emergency medical information in USA, call (877) 231-2615 or call collect 0 (952) 853-1713

Inactive ingredients water (aqua), isopropyl alcohol, propylene glycol, CI 16035 (FD&C Red 40), CI 42090 (FD&C Blue 1)

Questions? Call **1-800-529-5458**

Principal display panel and representative container

NDC 63146-123-10

Foaming Hand Sanitizer

KEEP OUT OF REACH OF CHILDREN

FOR INSTITUTIONAL USE ONLY

Benzalkonium chloride 0.1%

Net Contents:

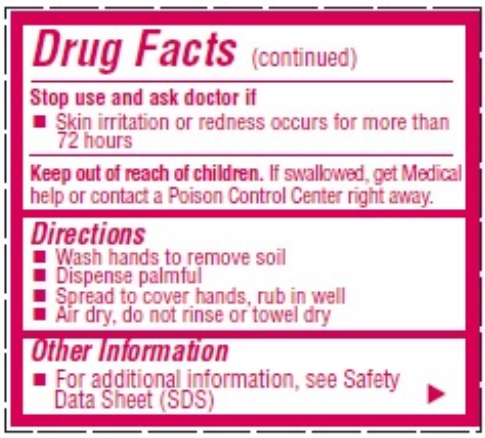
42 US fl oz (1250 ml)

FRSUSA 764026/8001/1020

Distributed by:

Kay Chemical Company · 8300 Capital Drive

Greensboro, NC 27409-9790 USA Customer Service: (800) 529-5458



ECOLAB

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63146-123
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63146-123-10	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/01/2016	

Labeler - Kay Chemical Company (003237021)

Revised: 12/2022 Kay Chemical Company