EXP FOAMING HAND SANITIZER - benzalkonium chloride solution
Kay Chemical Company

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

- 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 or 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.

Direction

- wet hands thoroughly with product and allow to dry without wiping.
- do not use if hands are visibly dirty or greasy; wash hands with soap and water instead.

Other information

- for additional information, see Safety Data Sheet (SDS)
- EMERGENCY HEALTH INFORMATION: 1 877 231 2615. If located outside the United States and Canada, call collect 952 853 1713 (number is in the US).

Inactive ingredients water (aqua), isopropyl alcohol, propylene glycol, FD&C red 40, FD&C blue 1

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

Principal display panel and representative label

ECOLAB

NDC 63146-320-16

EXP Foaming

Hand Sanitizer

Active Ingredient: Benzalkonium chloride 0.1%

1200 mL

(40.6 US FL OZ) 1112848

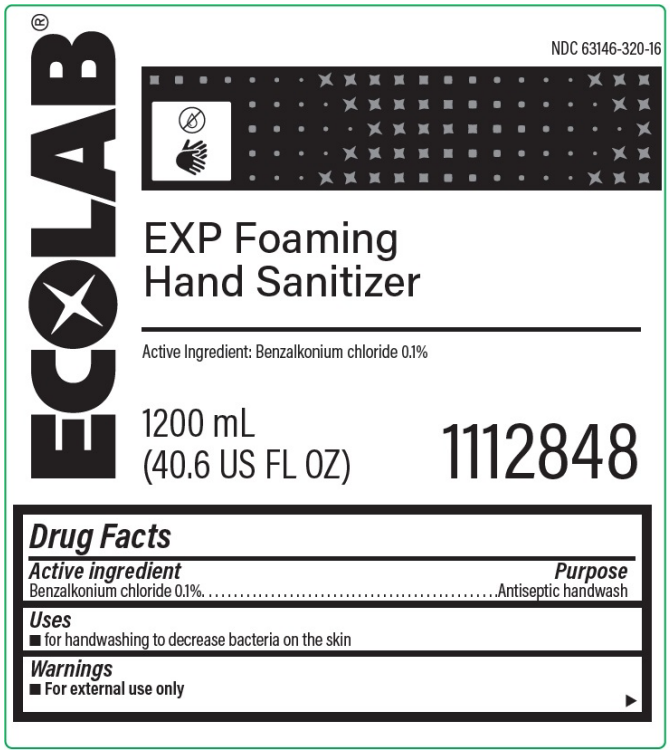
Kay Chemical Company · 8300 Capital Drive

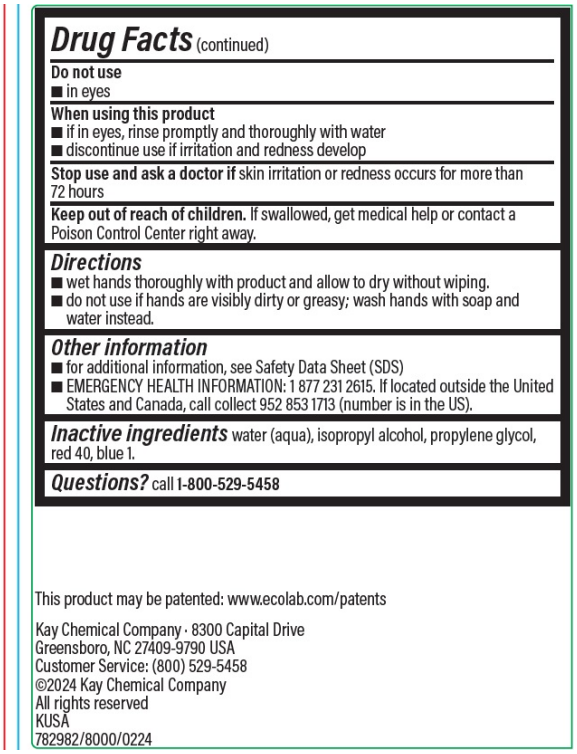
Greensboro, NC 27409-9790 USA

Customer Service: (800) 529-5458

KUSA

782982/8000/0224





EXP FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63146-320
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 mg in 100 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:63146-320-16	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/01/2024	

Labeler - Kay Chemical Company (003237021)

Revised: 7/2024 Kay Chemical Company