CHEMSTAR FOAMING HAND SANITIZER- 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

- for handwashing to decrease bacteria on skin

Warnings

For external use only

Do not use
- in eyes

When using this product
- if in eyes, rinse promptly and throughly with water
- discontinue use if irritation and redness develop

Stop use and ask doctor if
- skin irritation or redness persists 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 Safety Data Sheet (SDS)
- 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 label

CHEMSTAR

Foaming

Hand Sanitizer

Sanitizante de manos en espuma

KEEP OUT OF REACH OF CHILDREN

FOR INSTITUTIONAL USE ONLY

Benzalkonium chloride 0.1%

Net Contents: 42 US fl oz (1250 ml)

1111302

Distributed by:

Kay Chemical Company

8300 Capital Drive • Greensboro, NC 27409

Made in USA • CSUSA 778273/8000/0621

CHEMSTAR FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63146-316
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:63146-316-10	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/08/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/08/2010	

Labeler - Kay Chemical Company (003237021)

Revised: 11/2022 Kay Chemical Company