

CHEMSTAR FOAMING HAND SANITIZER- benzalkonium chloride solution
Ecolab Inc.

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 thoroughly 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

Drug Facts (continued)
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Drug Facts (continued)
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CHEMSTAR FOAMING HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-689
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:47593-689-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/18/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/18/2025	

Labeler - Ecolab Inc. (006154611)

Revised: 12/2025

Ecolab Inc.