CLEAN CHOICE FOODSERVICE ANTIBACTERIAL FOAM- benzalkonium chloride solution
SC Johnson Professional USA, Inc.

Drug Facts

Active ingredient

Benzalkonium Chloride, 0.13%

Purpose

Antibacterial

Uses

for hand washing to reduce bacteria on the skin

Warnings

For external use only

When using this product

avoid contact with eyes. In case of eye contact, flush with water.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

apply to dry hands

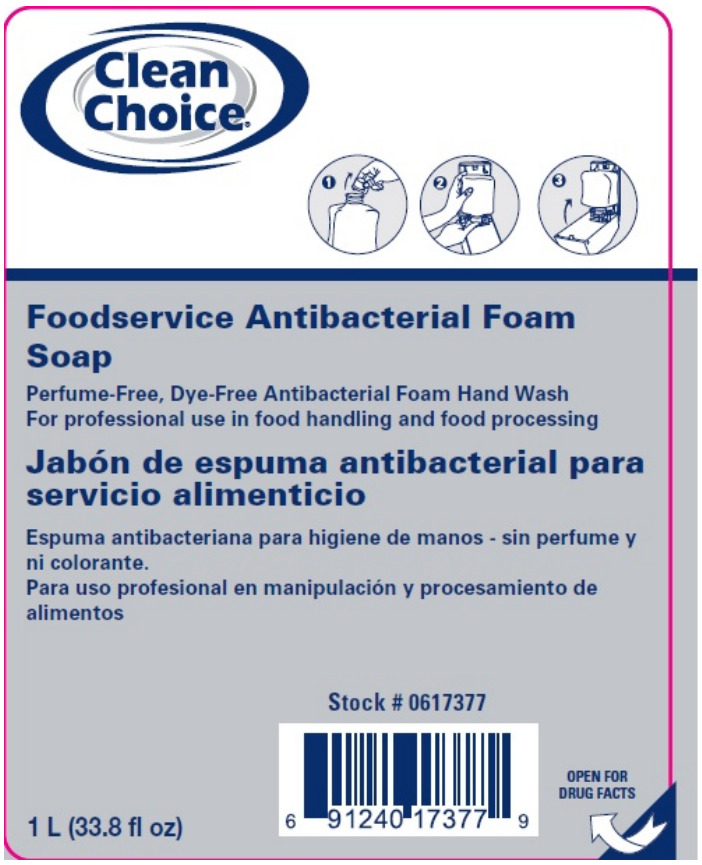
add water

lather hands for 30 seconds

rinse and dry hands thoroughly

Inactive ingredients

Aqua (Water), Hexylene Glycol, Coco-Glucoside, Cocamidopropyl PG-Dimonium Chloride Phosphate, Laurtrimonium Chloride, PEG-6 Cocamide, Citric Acid



Clean Choice

Foodservice Antibacterial Foam
Soap
Perfume-Free, Dye-Free Antibacterial Foam Hand Wash
For professional use in food handling and food processing
Jabón de espuma antibacterial para
servicio alimenticio
Espuma antibacteriana para higiene de manos - sin perfume y
ni colorante.
Para uso profesional en manipulación y procesamiento de
alimentos
Stock # 0617377
1 L (33.8 fl oz)
OPEN FOR
DRUG FACTS

CLEAN CHOICE FOODSERVICE ANTIBACTERIAL FOAM

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-025
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
COCO-GLUCOSIDE (UNII: ICS790225B)	
COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
PEG-6 COCAMIDE (UNII: YZ6NLA4O1E)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-025-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2018	
2	NDC:11084-025-66	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/17/2023	12/31/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	505G(a)(3)	10/01/2018	12/31/2026

Labeler - SC Johnson Professional USA, Inc. (607378015)

Establishment

Name	Address	ID/FEI	Business Operations
APEX International, Inc.		015226132	manufacture(11084-025)

Revised: 1/2025 SC Johnson Professional USA, Inc.