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

Drug Facts

Active ingredient

BENZALKONIUM CHLORIDE, 0.13%

Purpose

Antibacterial

Uses

for handwashing 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.

Stop use and ask a doctor if

irritation and redness develop and persist for more than 3 days.

Keep out of reach of children.

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

Directions

apply foaming cleanser to dry hands
rub hands together to spread lather
wash for 15-20 seconds
rinse & dry hands thoroughly

Inactive ingredients

AQUA (WATER), GLYCERIN, LAURAMINE OXIDE, BUTYLENE GLYCOL, LACTIC ACID, SALICYLIC ACID, PARFUM (FRAGRANCE), GREEN 5 (CI 61570), YELLOW 5 (CI 19140).

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TOTO
Antibacterial Foam Soap
✓ Cleans hands!
✓ Kills germs!
✓ Moisturizes!
✓ Triclosan-free!
One Gallon (3.78 Liters)
TSFG1
DCN9290
R00
Product Features
A high-quality, softly-perfumed, green foam soap that is 99.99% effective against common bacteria. This product contains moisturizers and emollients for the skin making it ideal for frequent use environments.
Manufactured for:
TOTO USA, Inc.
1155 Southern Road
Morrow, GA 30260
888-295-8134
Made in the USA

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ANTIBACTERIAL FOAM

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-021
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 g in 1 L

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	

FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11084-021-05	3.78 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/01/2017	
2	NDC:11084-021-42	18.93 L in 1 PAIL; Type 0: Not a Combination Product	08/01/2017	12/31/2024
3	NDC:11084-021-55	208.20 L in 1 DRUM; Type 0: Not a Combination Product	08/01/2017	12/31/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	505G(a)(3)	08/01/2017	

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

Establishment

Name	Address	ID/FEI	Business Operations
APEX International, Inc.		015226132	manufacture(11084-021)

Revised: 12/2024 SC Johnson Professional USA, Inc.