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

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, 0.13%

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

PRINCIPAL DISPLAY PANEL - 1 L Bottle Label

SCJ PROFESSIONAL
A Family Company™

NDC 11084-024-27

AgroBac™ PURE

ANTIMICROBIAL FOAM SOAP
Dye & Fragrance-Free E2 Antibacterial Foam Handwash.

NSF®
Nonfood Compounds
Program Listed E2
157532

1 L (33.8 fl oz)

SC Johnson Professional USA, Inc.
Charlotte, NC 28217
1-800-248-7190. www.scjp.com

Pat.www.scjp.com/patents

deb
SKIN CARE

1000002834/0320

AGROBAC PURE FOAM

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-024
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:11084-024-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2018	
2	NDC:11084-024-20	2000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2018	
3	NDC:11084-024-12	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2018	
4	NDC:11084-024-01	47 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/01/2018	07/19/2021
5	NDC:11084-024-66	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/17/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/01/2018	

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

Establishment

Name	Address	ID/FEI	Business Operations
APEX International, Inc.		015226132	manufacture(11084-024)

Revised: 2/2023 SC Johnson Professional USA, Inc.