REFRESH ANTIBAC FOAM- benzalkonium chloride soap
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

Antimicrobial

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).

PRINCIPAL DISPLAY PANEL - 1 L Bottle Label

SCJ Professional
A family company®

NDC 11084-010-66

Refresh™

ANTIBAC FOAM

Citrus Scent Antimicrobial Foam Handwash

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

CITRUS FRAGRANCE

deb
SKIN CARE

1 L
(33.8 fl oz)

Made in Canada
L-1467 R0

SCJ PROFESSIONAL
A family company®

NDC 11084-010-66

Refresh™ ANTIBAC FOAM

Citrus Scent Antimicrobial Foam Handwash

Drug Facts

Active ingredient ..... **Purpose**

Benzalkonium chloride 0.13% ..... Antimicrobial

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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REFRESH ANTIBAC FOAM

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 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:11084-010-40	0.4 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/12/2017	
2	NDC:11084-010-27	1 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/12/2017	
3	NDC:11084-010-12	1.2 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/12/2017	
4	NDC:11084-010-20	2 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/12/2017	
5	NDC:11084-010-05	3.78 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/12/2017	
6	NDC:11084-010-01	0.047 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/27/2019	
7	NDC:11084-010-55	208.2 L in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK; Type 0: Not a Combination Product	02/27/2019	
8	NDC:11084-010-96	0.296 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/01/2019	
9	NDC:11084-010-66	1 L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/31/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/12/2017	

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

Establishment

Name	Address	ID/FEI	Business Operations
APEX International, Inc.		015226132	manufacture(11084-010)

Establishment

Name	Address	ID/FEI	Business Operations
SC Johnson Professional CA Inc.		203765300	MANUFACTURE(11084-010)

Revised: 12/2022 SC Johnson Professional USA, Inc.