APPEAL ANTIBACTERIAL HAND WASH- 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 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.

Keep out of reach of children.

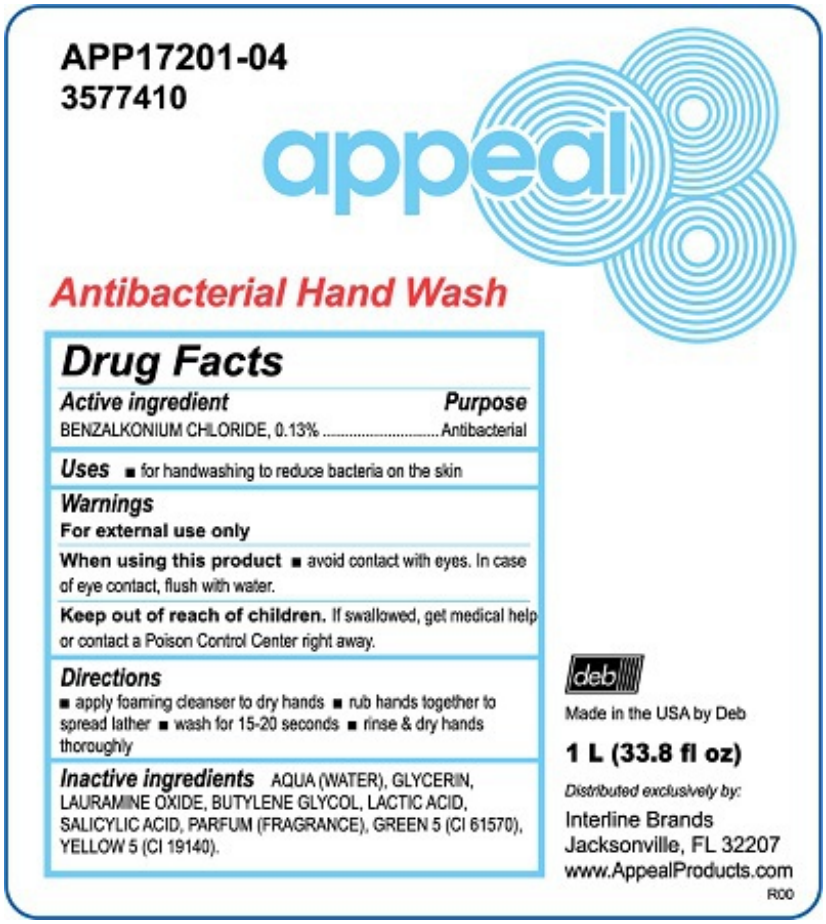
If swallowed, get medical help or contact a Poison Control Center right away.

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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Antibacterial Hand Wash

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Made in the USA by Deb

1 L (33.8 fl oz)

Distributed exclusively by:

Interline Brands

Jacksonville, FL 32207

www.AppealProducts.com

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APPEAL ANTIBACTERIAL HAND WASH

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11084-015
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 100 mL

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:11084-015-27	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/12/2017	
2	NDC:11084-015-66	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	06/12/2017	

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

Establishment

Name	Address	ID/FEI	Business Operations
APEX International, Inc.		015226132	MANUFACTURE(11084-015)

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