

AFIA FOAMING E2 SANITIZING HAND CLEANER- benzalkonium chloride soap
National Chemical Laboratories, 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.

Listing of Afia Foaming E2 Sanitizing Hand Cleaner

Drug Facts

Active Ingredient. Purpose

Benzalkonium Chloride 0.15%.....Antimicrobial hand Cleaner

Drug Facts		 7 5 2 6 1 0 7 0 4 4 8 8	National Chemical Laboratories, Inc. 401 N. 10th Street Philadelphia, PA 19123
Active Ingredient	Purpose		
Benzalkonium Chloride 0.15%.....	Antimicrobial Hand Cleaner		
Uses • For hand-washing to decrease bacteria on the skin. Helps prevent cross contamination by hand contact.			
Warnings			
For external use only.			
When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.			
Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.			
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.			
Directions • Wet hands and wrist with water. Pump 2 strokes of foam into palm of hand. • Rub thoroughly over all surfaces of both hands and wrist, include between fingers and under cuticles, for 60 seconds. • Rinse hands completely and dry thoroughly.			
Other Information • Do not contaminate potable water, food or feed, by use or storage or disposal.			
Inactive ingredients Water / Cetrimonium Chloride / Coco Glucoside / Dihydroxyethyl Cocamine Oxide / Laurtrimonium Chloride / Glycereth-17 Cocoate / Cocamidopropyl Hydroxysultaine / Hydroxyethylcellulose / Citric Acid			
			
Foaming E2 Sanitizing Hand Cleaner			
NDC 71023-448-57 LB1000-0001-0448-01i		1000mL (33.8 fl. oz.) Product #0448	

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afia™
skin care solutions

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World Class Cleaning Solutions®



0448-57

FOAMING E2 SANITIZING
HAND CLEANER



6/1 LITER BAGS

National Chemical Laboratories, Inc.
Philadelphia, PA 19123 USA • (800) 628-2436 • www.nclonline.com

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benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71023-448
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	
HYDROXYETHYL CELLULOSE (5000 MPAS AT 1%) (UNII: X70SE62ZAR)	
GLYCERETH-17 COCOATE (UNII: 3057VPT0KC)	

WATER (UNII: 059QF0KO0R)	
COCO GLUCOSIDE (UNII: ICS790225B)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71023-448-57	1000 mL in 1 BAG; Type 0: Not a Combination Product	01/18/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/18/2017	

Labeler - National Chemical Laboratories, Inc. (002289619)

Registrant - National Chemical Laboratories, Inc. (002289619)

Establishment

Name	Address	ID/FEI	Business Operations
National Chemical Laboratories, Inc.		002289619	manufacture(71023-448)

Revised: 1/2017

National Chemical Laboratories, Inc.