AFIA ALCOHOL-FREE FOAMING HAND SANITIZER- 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 Alcohol-Free Foaming Hand Sanitizer in multiple packages

Drug Facts

Active Ingredient. Purpose

Benzalkonium Chloride 0.1%.....Antimicrobial

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Active Ingredient. ***Purpose***
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7 52610 70445 7

National Chemical Laboratories, Inc.
401 N. 10th Street
Philadelphia, PA 19123

Uses • For hand sanitizing to decrease bacteria on the skin • Recommended for repeated use

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

• Pump a small amount of foam into palm of hand • Rub thoroughly over all surfaces of both hands • Rub hands together briskly until dry

Inactive ingredients

Water, dihydroxypropyl PEG-5 linoleammonium chloride, glycereth-2 cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide, fragrance

Alcohol-Free Foaming Hand Sanitizer

afia™ skin care solutions

NDC 71023-445-57
LB1000-0001-0445-01i

1000mL (33.8 fl. oz.)
Product #0445

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Alcohol-Free Foaming
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afia


NCL
World Class Cleaning Solutions
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0445-57
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FOAMING HAND SANITIZER
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AFIA ALCOHOL-FREE FOAMING HAND SANITIZER

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71023-445
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
GLYCERETH-2 COCOATE (UNII: JWM00VS7HC)	
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71023-445-57	1000 mL in 1 BAG; Type 0: Not a Combination Product	10/24/2016	
2	NDC:71023-445-29	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/01/2020	
3	NDC:71023-445-32	208450 mL in 1 DRUM; Type 0: Not a Combination Product	03/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/24/2016	

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(71023-445)

Revised: 10/2022 National Chemical Laboratories, Inc.