FOAMING HAND SANITIZER- foaming hand sanitizer liquid
National Chemical Laboratories Of PA., Inc.

AFIA F207F NCL

Active Ingredient - Benzalkonium Chloride 0.13% w/w.

Purpose - Antibacterial Agent

Uses - Hand sanitizer to help reduce bacteria on the skin that could cause disease.

Warnings

For external use only.

Avoid contact with eyes. If contact occurs, flush wyes with water.

Stop use and ask a doctor if, in rare instances, redness or irritation develops and persists for more than 72 hours.

Keep out of reach of children. If swallowed, contact a phsician or poison control.

Directions

Apply a small amount to palm. Briskly rub, covering hands with product until dry.

Inactive ingredients

Water, Propylene Glycol, Cocamidopropyl Betaine, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate (Vitamin E), PEG-7 Glyceryl Cocoate, Fragrance/Parfum, Phenoxyethanol, Tetrasodium EDTA.

Drug Facts

Active Ingredient — **Purpose**
Benzalkonium Chloride 0.13% w/w..Antibacterial Agent

7 52610 70445 7

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

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LAM1473S

Alcohol-Free Foaming Hand Sanitizer

afia™
skin care solutions

LB1000-0001-0445-03

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

FOAMING HAND SANITIZER

foaming hand sanitizer liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71023-207
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FRAGRANCE LINEN ORC1900779 (UNII: 658A7B6MY1)	
PEG-7 GLYCERYL COCOATE (UNII: VNX7251543)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71023-207-41	1000 mL in 1 BAG; Type 0: Not a Combination Product	06/18/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/18/2024	

Labeler - National Chemical Laboratories Of PA., Inc. (002289619)

Registrant - Kutol Products Company (004236139)

Establishment

Name	Address	ID/FEI	Business Operations
Kutol Products Company		004236139	manufacture(71023-207)

Revised: 10/2024 National Chemical Laboratories Of PA., Inc.