FOAMING HAND- benzalkonium chloride lotion Cintas Corp

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.

Sanitizing Foaming Hand Soap 224.000/224AA

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Uses

for handwashing to decrese bacteria on the skin

Warnings

for external use only: hands only

When using this product

• avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor if

- irritation or redness develop
- condition persists for more than 72 hours

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

Other Information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finshes and plastics

Inactive ingredients

water, cocamidopropyl betaine, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, glycerin, citric acid, tetrasodium EDTA, sodium benzoate

Rear label text

Manufactured for Cintas Corp.

Mason, OH 45040

800-CINTAS1

NSF Nonfood Compounds Program

Product is NSF certified for use as a hand sanitizer in and around food processing areas.

CINTAS

800mL/27.0 FL OZ

principal display panel

HAND

SOAP

ANTIBACTERIAL

FOAM



FOAMING HAND

benzalkonium chloride lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:42961-224

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		
WATER (UNII: 059QF0KO0R)			
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)			
LAURAMIDOPROPYLAMINE OXIDE (UNII: 16KX160QTV)			

LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

F	Packaging			
#	# Item Code Package Description		Marketing Start Date	Marketing End Date
1	NDC:42961- 224-71	800 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/20/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/20/2020	
final	PS. 1555 .	07,23,232	

Labeler - Cintas Corp (056481716)

Registrant - Vi-Jon, LLC (790752542)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(42961-224)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(42961-224)

Revised: 2/2023 Cintas Corp