FOAMING HAND SANITIZER REFILL MOISTURIZING ALOE- benzalkonium chloride liquid Bocks 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.

Foaming Hand Sanitizer Refill (Moisturizing Aloe)

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- hand sanitizer that helps decrease bacteria on the skin
- recommend for repeated use

Warnings

For external use only, do not use in ears, mouth, or eyes.

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

Children should be supervised when using this product.

Directions

- pour liquid into a foaming dispenser
- pump a small amount of foam into palm of hand
- spread thoroughly on both hands
- rub hands together until dry

Inactive Ingredients

Aloe Juice, Citric Acid, Dimonium Chloride Cocoamidopropyl Hydroxyethyldimonium Chloride, Hydroxypropyl Bis-Hydroxyethyldimonium Chloride, Rose Ether, Sodium Citrate, Water

Package Labeling: 473ml



Package Labeling: 946ml



FOAMING HAND SANITIZER REFILL MOISTURIZING ALOE

benzalkonium chloride liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81527-001	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
CITRIC ACID MONO HYDRATE (UNII: 2968 PHW8 QP)		
HYDROXYPROPYL BIS-HYDROXYETHYLDIMONIUM CHLORIDE (UNII: 1EO3O1X7HG)		
SO DIUM CITRATE (UNII: 1Q73Q2JULR)		
WATER (UNII: 059QF0KO0R)		

I	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:81527-001-16	473 mL in 1 CARTON; Type 0: Not a Combination Product	02/01/2021	
	NDC:81527-001-32	946 mL in 1 CARTON; Type 0: Not a Combination Product	02/01/2021	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	02/01/2021		

Labeler - Bocks Inc (082200343)

Revised: 2/2021 Bocks Inc