# FOAMING HAND SANITIZER REFILL FRAGRANCE FREE- 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.

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#### Foaming Hand Sanitizer Refill (Fragrance Free)

#### **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

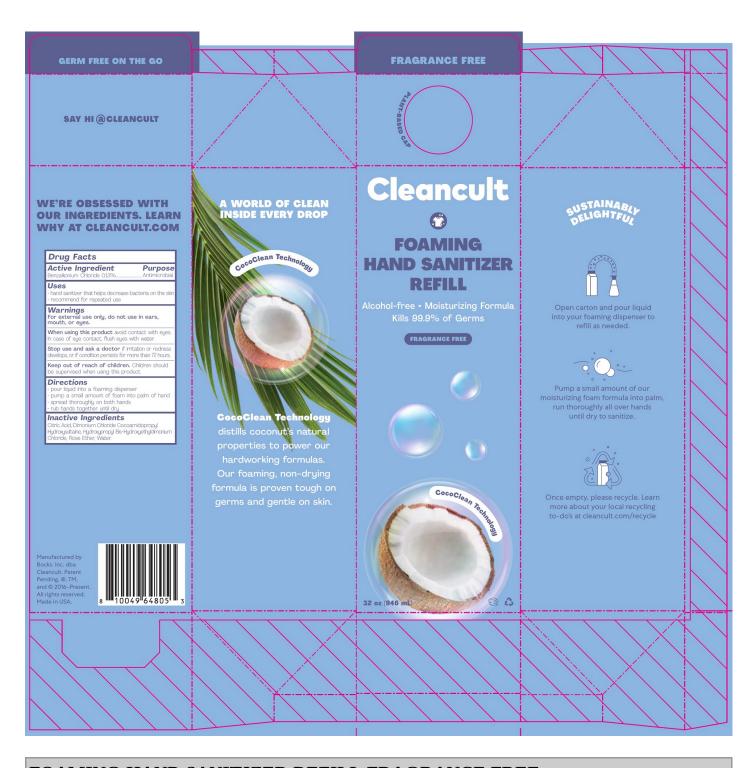
#### **Inactive Ingredients**

Citric Acid, Dimonium Chloride Cocoamidopropyl Hydroxysultaine, Hydroxypropyl Bis-Hydroxyethyldimonium Chloride, Rose Ether, Water.

### Package Labeling: 473ml



Package Labeling: 946ml



#### FOAMING HAND SANITIZER REFILL FRAGRANCE FREE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81527-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM -	BENZALKONIUM	1.3 mg

UNII:7N6JUD5X6Y)	CHLORIDE	in 1 mL
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Inactive Ingredients		
Ingredient Name	Strength	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)		
HYDRO XYPRO PYL BIS-HYDRO XYETHYLDIMO NIUM CHLO RIDE (UNII: 1EO3O1X7HG)		
WATER (UNII: 059QF0KO0R)		

l	Packaging			
l	# Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
l	1 NDC:81527-000-16	473 mL in 1 CARTON; Type 0: Not a Combination Product	02/01/2021	
	2 NDC:81527-000-32	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