# HEALTHY ACCENTS ANTIBACTERIAL FOAMING HAND- benzalkonium chloride soap DZA Brands LLC

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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### **Drug Facts**

#### **Active ingredient**

Benzalkonium Chloride 0.13%

#### Purpose

Antibacterial

#### Uses

• For washing to decrease bacteria on the skin.

### **Warnings**

For external use only.

#### When using this product

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

Stop using this product and ask doctor if

• irritation or redness develops and lasts.

Keep out of reach of children.

• In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

#### **Directions**

• Pumpm onto dry hands, work into a lther. Rinse thoroughly.

#### **Questions? Comments?**

1-866-322-2439

#### **Inactive ingredients**

Water (Aqua), Cocamidopropyl Betaine, Polysorbate 20, Fragrance (Parfum), Glycerin, Decyl Glucoside, Hydroxyethylcellulose, Aloe Barbadensis Leaf Juice, Camellia Sinensis Leaf Extract, Polyquaternium-7, Tetrasodium EDTA, Sodium Citrate, Citric Acid, Benzophenone-4, Methylchloroisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700), Yellow 5 (CI 19140).

## **Label Copy**





### HEALTHY ACCENTS ANTIBACTERIAL FOAMING HAND

benzalkonium chloride soap

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:55316-196

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name

Basis of Strength

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM UNII:7N6JUD5X6Y)

BENZALKONIUM
CHLORIDE

1.3 mg
in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3 O 11 KX)				
POLYSORBATE 20 (UNII: 7T1F30V5YH)				
GLYCERIN (UNII: PDC6A3C0OX)				
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)				
HYDROXYETHYL CELLULOSE (5000 CPS AT 1%) (UNII: X70SE62ZAR)				
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
GREEN TEA LEAF (UNII: W2ZU1RY8B0)				

POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
EDETATE SO DIUM (UNII: MP1J8420 LU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
CITRIC ACID MONO HYDRATE (UNII: 2968 PHW8 QP)	
SULISOBENZONE (UNII: 1W6L629B4K)	
METHYLCHLORO ISO THIAZO LINO NE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)	
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

	Packaging					
	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
ı	1 NDC:55316-196- 07	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/06/2017			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333E	06/06/2017				

## Labeler - DZA Brands LLC (090322194)

## **Registrant** - Apollo Health and Beauty Care Inc. (201901209)

Establishment						
Name	Address	ID/FEI	Business Operations			
Apollo Health and Beauty Care Inc.		201901209	manufacture(55316-196)			

Revised: 6/2017 DZA Brands LLC