P.O.V. SPRINGTIME RENEWAL ANTIBACTERIAL FOAMING HAND- benzalkonium chloride soap
Apollo Health and Beauty Care 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.

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

helps eliminate bacteria on hands.

Warnings

For external use only.

When using this product

avoid contact with eyes. In case of contact, rinse thoroughly with water.

Stop use and ask a doctor if

irritation and 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

- apply onto dry hands.
- lather and rinse thoroughly.

Other information

store at room temperature.

Inactive ingredients

Water (Aqua), Cocamidopropyl Betaine, Polysorbate 20, Fragrance (Parfum), Glycerin, Decyl Glucoside Hydroxyethylcelluose, Aloe Barbadensis Leaf Juice, Camellia Sinensis Leaf Extract, Propylen Glycol, Polyquaternium-7, Tetrasodium EDTA, Sodium Citrate, Citric Acid, Benzophenone-4, Methylchloroisothiazolinone, Methylisothiazolinone, Blue 1 (CI 42090), Ext. Violet 2 (CI 60730).

Questions or comments?

1-866-695-3030

Label Copy

P.O.V. SPRINGTIME RENEWAL ANTIBACTERIAL FOAMING HAND

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63148-198
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: 5OCF3O11KX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
GLYCERIN (UNII: PDC6A3C0OX)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
HYDROXYETHYL CELLULOSE (5000 CPS AT 1%) (UNII: X70SE62ZAR)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SULISOBENZONE (UNII: 1W6L629B4K)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63148-198-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/10/2017	
2	NDC:63148-198-32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/10/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/10/2017	

Labeler - Apollo Health and Beauty Care Inc. (201901209)

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

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(63148-198)

Revised: 11/2017 Apollo Health and Beauty Care Inc.