DAYLOGIC FRESH PEAR FOAMING- benzalkonium chloride soap Rite Aid Corporation

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

for hand washing to decrease bacteria on the skin

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 develop and lasts.

Keep out of reach of children.

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

Directions

- pump onto dry hands
- work into a rich foamy lather, rinse thoroughly and dry

Other information

store at room temperature

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, Yellow 10 (CI 47005), Green 3 (CI 42053)

Label Copy



DAYLOGIC FRESH PEAR FOAMING

benzalkonium chloride soap

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11822-1090

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: 7T1F30 V5YH)		
GLYCERIN (UNII: PDC6A3C0OX)		
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)		
HYPROMELLOSES (UNII: 3NXW29 V3WO)		
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: MP1J8 420 LU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
SULISOBENZONE (UNII: 1W6L629B4K)	
METHYLCHLORO ISO THIAZO LINO NE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	

ı	P	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1	NDC:11822-1090- 8	222 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/09/2017			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333E	05/09/2017				

Labeler - Rite Aid Corporation (014578892)

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

Establishment				
Name	Address	ID/FEI	Business Operations	
Apollo Health and Beauty Care Inc.		201901209	manufacture(11822-1090)	

Revised: 5/2017 Rite Aid Corporation