

TOPCARE ANTIBACTERIAL FOAMING HAND FRESH PEAR- benzalkonium chloride liquid
TopCo Associates 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.

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

helps eliminate 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 using this product and ask a 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

Use only to refill a foaming hand soap pump bottle. From pump bottle, apply onto dry hands work into a lather and rinse thoroughly.

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, Polyquatonium-7, Xanthan Gum, Tetrasodium EDTA, Sodium Citrate, Citric Acid, Methylchloroisothiazolinone, Methylisothiazolinone, Yellow 10 (CI 47005), Green 3 (CI 42053).

Label Copy

TopCare®

ANTIBACTERIAL foaming hand soap

gentle enough
for every day

FRESH PEAR

32 FL OZ (1 QT) 946 mL

06-19709

TopCare®

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Directions use only to refill a Foaming Hand Soap from pump bottle, apply onto dry hands work into a lather and rinse thoroughly.

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DISTRIBUTED BY
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MADE IN CANADA 06-23148



TOPCARE ANTIBACTERIAL FOAMING HAND FRESH PEAR

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:36800-109
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)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 160000 MW) (UNII: 0L414VCS5Y)	
XANTHAN GUM (UNII: TTV12P4NEE)	
EDETATE SODIUM (UNII: MPIJ8420LU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-109-33	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/28/2017	
2	NDC:36800-109-07	222 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/28/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/28/2017	

Labeler - TopCo Associates LLC (006935977)

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

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(36800-109)