# TOPCARE EVERYDAY SPRING FRESH BODY WASH- 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.

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## **Drug Facts**

# **Active ingredient**

Benzalkonium Chloride 0.13%

# Purpose

Antibacterial

#### Use

For washing to decrease bacteria on the skin.

### Warnings

For external use only.

### When using this product

avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop using this product 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**

Squeeze a small amount onto a wet wash cloth, sponge, pouf or hands and apply to body. Work into a rich lather and rinse off.

#### Questions or comments?

1-888-423-0139

#### **Inactive ingredients**

Water (Aqua), Cocamidopropyl Betaine, Decyl Glucoside, Hydroxyethylcellulose, Glycerin, Poloxamer 124, Fragrance (Parfum), Aloe Barbadensis Leaf Juice, Polyquaternium-7, Tetrasodium EDTA, Citric Acid, Propylene Glycol, Sodium Citrate, Methylchloroisothiazolinone, Methylisothiazolinone, Blue 1 (CI 42090), Ext. Violet 2 (CI 60730).

#### **Label Copy**



### TOPCARE EVERYDAY SPRING FRESH BODY WASH

benzalkonium chloride liquid

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:36800-895
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
<b>BENZALKONIUM CHLORIDE</b> (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)		
<b>DECYL GLUCOSIDE</b> (UNII: Z17H97EA6Y)		
HYDROXYETHYL CELLULOSE (5000 CPS AT 1%) (UNII: X70SE62ZAR)		
GLYCERIN (UNII: PDC6A3C0OX)		
POLOXAMER 124 (UNII: 1S66E28KXA)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)		
EDETATE SO DIUM (UNII: MP1J8420LU)		
CITRIC ACID MONOHYDRATE (UNII: 2968 PHW8 QP)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
SODIUM CITRATE (UNII: 1Q73Q2JULR)		
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)		
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)		
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:36800-895- 18	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/30/2019	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	10/30/2019		

# 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(36800-895)

Revised: 10/2019 Topco Associates LLC