

ATERRA7 ANTIBACTERIAL FOAMING HAND- benzalkonium chloride soap
B4 Ventures LLC

Aterra7 Antibacterial Foaming Hand Soap

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Skin Antimicrobial

Use

reduces amount of bacteria on hands

Warnings

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Apply a palmful to hands and forearms.
- Scrub thoroughly for at least 15 seconds.
- Rinse completely and dry.

Inactive Ingredients

Water, Lauramine Oxide, Glycerin, PEG-120 Methyl Glucose Dioleate, DMDM Hydantoin, Citric Acid, Fragrance

Leaves Hands
Feeling Soft &
Refreshed



Aterra[®]7

NDC 68306-115-03

LBL 1666-2.0

Antibacterial
FOAMING HAND SOAP

Triclosan-Free

Leaves Hands Feeling Soft & Refreshed

33.8 fl. oz. (1 qt. 1.8 fl. oz.) 1000 mL

Drug Facts

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Item #: 12007-9410
LBL1667-2.1

Not tested on animals



Made in USA

Manufactured for B4 Brands, Lisbon, Iowa 52253

ATERRA7 ANTIBACTERIAL FOAMING HAND

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68306-115
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Product Characteristics

Color	white (water white, colorless - dispensed as white foam)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68306-115-01	530 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/26/2018	
2	NDC:68306-115-02	4 in 1 BOX	09/26/2018	
2		1000 mL in 1 BAG; Type 0: Not a Combination Product		
3	NDC:68306-115-03	4 in 1 BOX	09/20/2019	
3		1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/26/2018	

