AVANT ALCOHOL-FREE INSTANT HAND SANITIZER- benzalkonium chloride solution
B4 Ventures LLC

Avant Alcohol-Free Foaming Instant Hand Sanitizer

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antimicrobial

Use

For hand sanitizing to decrease bacteria on the skin

Warnings

For external use only.

Do not use in the eyes. In case of eye contact, flush thoroughly with water and seek medical attention.

Stop use and ask doctor if irritation and redness develop and 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

- Dispense an adequate amount in your palm to cover all surfaces of hands completely.
- Rub hands together until dry.
- Supervise children in the use of this product.

Inactive Ingredients

Water, Glycerin, Propylene Glycol, Phenoxyethanol, Lauramine Oxide, Panthenol, Citric Acid

Questions or Comments?

1-888-667-6066 or www.b4brands.com

AVANT ALCOHOL-FREE INSTANT HAND SANITIZER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68306-112
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 g in 1000 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
PANTHENOL (UNII: WV9CM0O67Z)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68306-	530 mL in 1 BOTTLE, PUMP; Type 0: Not a	01/01/2012	

1	112-01	Combination Product	01/01/2012	
2	NDC:68306-112-02	1000 mL in 1 POUCH; Type 0: Not a Combination Product	01/01/2012	
3	NDC:68306-112-03	2000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/01/2012	09/26/2024
4	NDC:68306-112-04	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	01/01/2012	
5	NDC:68306-112-05	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/27/2023	
6	NDC:68306-112-06	208198 mL in 1 DRUM; Type 0: Not a Combination Product	05/09/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/2012	

Labeler - B4 Ventures LLC (133582853)

Revised: 8/2024 B4 Ventures LLC