AVANT FOAMING HAND SANITIZER- alcohol liquid
B4 Ventures LLC

B4 Brands: Avant Foaming Hand Sanitizer - Fragrance Free

Active Ingredient

Ethanol 62% v/v

Purpose

Antimicrobial

Uses

- Hand sanitizer to help reduce bacteria on the skin
- Recommended for repeated use

Warnings

For external use only.

Flammable, keep away from fire or flame.

When using this product do not use in or near eyes. In case of eye contact, flush thoroughly with water.

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.

Other Information

- Do not store above 100 ºF (38 ºC)
- May discolor some fabrics

Inactive Ingredients

Water, Acrylates/Perfluorohexylethyl Methacrylate Copolymer, Perfluorohexylethyl

Alcohol, Glycerin, Aloe Barbadensis Leaf Juice

Questions or comments?

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

AVANT FOAMING HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68306-109
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
2-(PERFLUOROHEXYL)ETHANOL (UNII: G2R5YO5N3V)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BUTYL ACRYLATE/METHYL METHACRYLATE/PERFLUOROHEXYLETHYL METHACRYLATE COPOLYMER (SALUS AF) (UNII: HLB263IJK9)	
GLYCERIN (UNII: PDC6A3C0OX)	

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:68306-109-01	530 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/08/2010	
2	NDC:68306-109-02	1000 mL in 1 POUCH; Type 0: Not a Combination Product	12/08/2010	
3	NDC:68306-109-03	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	05/01/2012	
4	NDC:68306-109-05	1040988 mL in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK; Type 0: Not a Combination Product	10/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/08/2010	

Labeler - B4 Ventures LLC (133582853)

Revised: 10/2024 B4 Ventures LLC