ARTEMIS ALCOHOL-FREE ANTISEPTIC FOAM HAND SANITIZER- benzalkonium chloride liquid
Artemis Bio-Solutions, Inc.

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.

ARTEMIS ALCOHOL-FREE Antiseptic Foam Hand Sanitizer

Drug Facts

Active Ingredients

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

- For external use only.
- Do not use in eyes. If contact occurs, flush eyes with water.
- Stop use and ask a doctor if irritation or redness develops. If condition persists for more than 72 hours, consult a doctor.
- Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Pump a small amount of foam into palm of hand.
- Wet hands thoroughly with product and allow to dry without wiping.
- Rub hands together briskly until dry.

Inactive Ingredients

Water, dihydroxypropyl PEG-5 linoleammonium chloride, glycereth-2cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide, fragrance

ARTEMIS BIOSOLUTIONS

Manufactured by Artemis BioSolutions.

960 N. Industrial Dr., Ste.4 Elmhurst, IL 60126.

Made in USA.

- NO RINSE
- MOISTURIZES
- LEAVES SKIN SOFT

ARTEMIS
Many threats. One solution

ALCOHOL-FREE
Antiseptic Foam Hand Sanitizer

Eliminates 99.999%
OF MOST COMMON GERMS THAT MAY CAUSE ILLNESS

Environmentally Friendly
1.7 floz (50ml)

ARTEMIS ALCOHOL-FREE ANTISEPTIC FOAM HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49765-310
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 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
DIHYDROXYPROPYL PEG-5 LINOLEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)	
GLYCERETH-2 COCOATE (UNII: JWM00VS7HC)	
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)	
DIHYDROXYETHYL COCAMINE OXIDE (UNII: 8AR51R3BL5)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:49765-310-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/24/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/24/2011	

Labeler - Artemis Bio-Solutions, Inc. (116934417)

Establishment

Name	Address	ID/FEI	Business Operations
Artemis Bio-Solutions, Inc.		116934417	manufacture(49765-310)

Revised: 1/2020 Artemis Bio-Solutions, Inc.