ZALIEX ALCOHOL-FREE FOAMING HAND SANITIZER - benzalkonium chloride liquid
SAS Healthcare 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.

Zaliex Alcohol-Free Foaming Hand Sanitizer

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Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antimicrobial

Uses:
For hand sanitizing, to eliminate harmful bacteria and germs on skin.
Use as part of your daily cleansing routine.
Recommended for repeated use.

Warnings:

For external use only.

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

Stop use and ask a doctor if irritation or redness develops, or if condition persists.

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
- Run thoroughly over all surfaces of both hands
- Rub hands together briskly until dry

Inactive Ingredients:

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

Manufactured for:

Zaliex

Suite 210, 4 Robert Speck Parkway,

Mississauga, Ontario, L4Z1S1, Canada

Zaliex Alcohol-Free Foaming Hand Sanitizer 1L (76452-004-00)

ZALIEX ALCOHOL-FREE FOAMING HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76452-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 mg in 0.001 L

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:76452-004-00	1 L in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/26/2011	

Labeler - SAS Healthcare Inc (248055696)

Establishment

Name	Address	ID/FEI	Business Operations
Artemis Bio-Solutions Inc.		963442541	manufacture(76452-004)

Revised: 9/2012 SAS Healthcare Inc