ZALIEX ALCOHOL-FREE ANTISEPTIC 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 Antiseptic Foam Hand Sanitizer

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Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses:

- For hand cleansing 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-2 cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide, fragrance

Manufactured for:

Zaliex

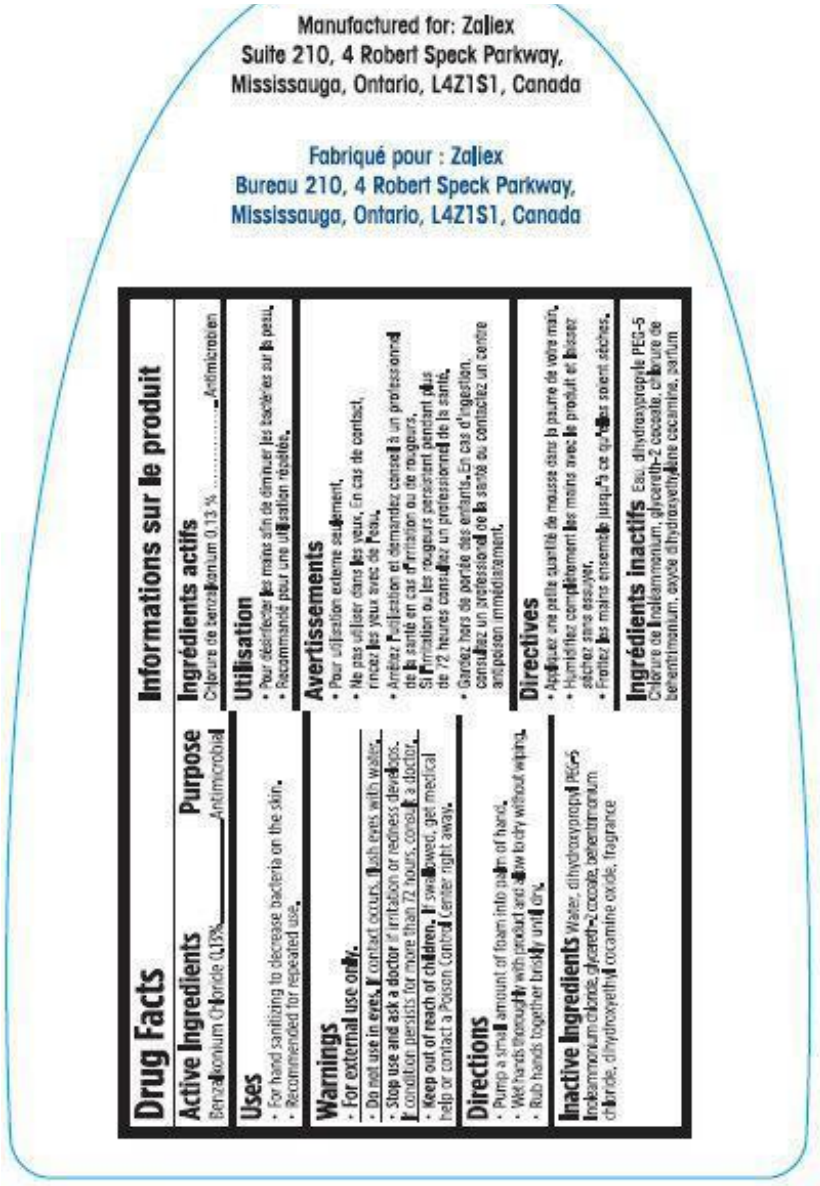
Suite 210, 4 Robert Speck Parkway,

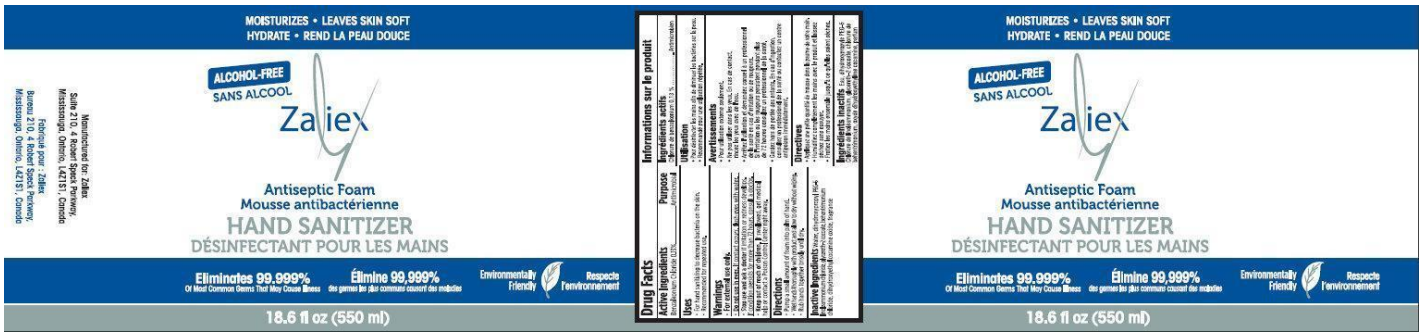
Mississauga, Ontario, L4Z1S1, Canada

Zaliex Alcohol-Free Antiseptic Foam Hand Sanitizer 250ml (76452-003-00) | Zaliex Alcohol-Free Antiseptic Foam Hand Sanitizer 550ml (76452-003-01)


ALCOHOL-FREE
SANS ALCOOL
Zaliex
Antiseptic Foam
Mousse antibactérienne
HAND SANITIZER
DÉSINFECTANT POUR LES MAINS
MOISTURIZES • LEAVES SKIN SOFT
HYDRATE • REND LA PEAU DOUCE
Eliminates 99.999%
Of Most Common Germs That May Cause Illness
Élimine 99,999%
des germes les plus communs causant des maladies
8.4 fl oz (250 ml)
Environmentally Friendly
Respecte l'environnement

2.375″





ZALIEX ALCOHOL-FREE ANTISEPTIC HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76452-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 mg in 1 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:76452-003-00	250 mL in 1 BOTTLE		
2	NDC:76452-003-01	550 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/13/2012	

Labeler - SAS Healthcare Inc (248055696)

Establishment

Name	Address	ID/FEI	Business Operations
Artemis Bio-Solutions Inc.		963442541	manufacture(76452-003)

Revised: 9/2012 SAS Healthcare Inc