

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

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

MOISTURIZES • LEAVES SKIN SOFT
HYDRATE • REND LA PEAU DOUCE

ALCOHOL-FREE
SANS ALCOOL



**FOAMING
HAND SANITIZER**
ASSAINISSEUR MOUSSANT
POUR LES MAINS

Eliminates harmful bacteria and germs on skin
Élimine les bactéries et les dangereux nocifs sur la peau

35.395 US fl. oz. (1 L)

DIN 02328240

Drug Facts	La Drogue Faits	Objectif	
Active Ingredients Benzalkonium Chloride 0.1%.....Antimicrobial	Ingrédients actifs Chlorure de benzalkonium 0,1%.....Antimicrobien		
Uses For hand sanitizing, to eliminate harmful bacteria and germs on skin. Use as part of your daily cleansing routine. Recommended for repeated use.	Utilisation Pour l'assainissement des mains en vue d'éliminer les bactéries et les dangereux nocifs sur la peau. Intégrez le produit dans votre routine de lavage quotidien. Recommandé pour une utilisation répétée.		
Warnings • For external use only. • When using this product avoid contact with eyes. In case of eye contact, flush eyes with water. Stop use and ask a physician if irritation or redness develops, or condition persists. • Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	Avertissements • Pour utilisation externe seulement. • Lorsque vous utilisez ce produit, ne res. le mettre en contact avec les yeux. En cas de contact avec les yeux, rincez les yeux à l'eau. Arrêtez l'utilisation et demandez conseil à un professionnel de la santé en cas d'irritation ou de rougeurs, ou si l'irritation ou les rougeurs persistent. • Gardez hors de portée des enfants. En cas d'ingestion, consultez un professionnel de la santé ou contactez un Centre Antipoison immédiatement.		
Directions • Pump a small amount of foam into palm of hand. • Rub thoroughly over all surfaces of both hands. • Rub hands together briskly until dry.	Directives • Appliquez une petite quantité de mousse dans la paume de votre main. • Frottez bien toute la surface des deux mains. • Frottez les mains ensemble jusqu'à ce qu'elles soient sèches.		
Inactive Ingredients Water, dihydroxypropyl PEG-5 linoleammonium chloride, glycereth-2 cocoate, behentrimonium chloride, dihydroxyethyl cocamine oxide, fragrance	Ingrédients inactifs Eau, dihydroxypropyle PEG-5 chlorure de linoléammonium, glycéth-2 coccoate, chlorure de behentrimonium, oxyde dihydroxyéthyle cocamine, parfum		



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Manufactured for Zalix • Suite 210, 4 Robert Speck Parkway,
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Fabrique pour Zalix • Bureau 210, 4 Robert Speck Parkway,
Mississauga, Ontario, L4Z1S1, Canada

ZALIX 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)				
DIHYDRO XYPROPYL PEG-5 LINO LEAMMONIUM CHLORIDE (UNII: 0Y0NQR2GH1)				
GLYCERETH-2 CO COATE (UNII: JWM00VS7HC)				
BEHENTRIMONIUM CHLORIDE (UNII: X7GNG3S47T)				
DIHYDRO XYETHYL CO CAMINE 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