SANITIZING HAND WIPES- benzalkonium chloride cloth
Landy International

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.

51706-907 Sanitizing wipes 0.13% Benzalkonium Chloride

Active Ingredient(s)

Benzalkonium Chloride 0.13%

Purpose

Antiseptic/Hand & Skin Sanitizer

Use

Sanitizing Hand Wipes to help decrease bacteria on the skin.
Recommended for single use.

Warnings

Do not freeze. For external use only.
Do not use in ears, eyes or mouth.
●Avoid contact with the eyes. In case of contact, flush eyes with water.
●Stop use and ask a doctor if redness or iritation
develop and persist for more than 72 hours.
●Keep out of reach of children.
●Children should be supervised when using this product.

Keep out of reach of children.

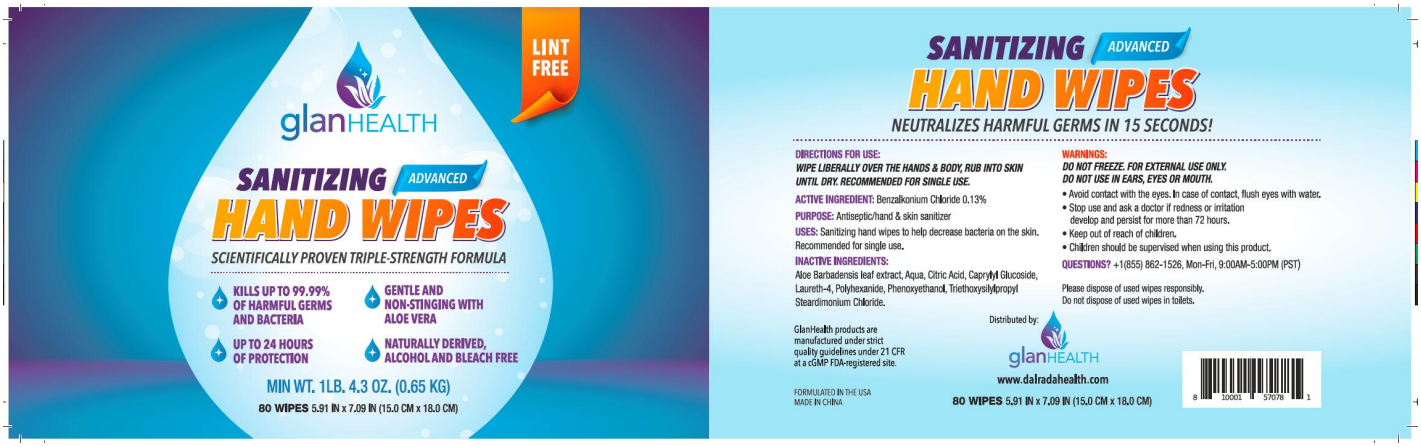
Directions

Wipe liberally over the hands & body, rub into skin until dry.
Recommended for single use.

Inactive ingredients

Aloe Barbadensis leaf extract, Aqua, Citric Acid, CaprylyI Glucoside, Laureth-4, Polyhexanide, Phenoxyethanol, Triethoxysilylpropyl Steardimonium Chloride.

Package Label - Principal Display Panel



SANITIZING HAND WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51706-907
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 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
TRIETHOXY(3-ISOCYANATOPROPYL)SILANE (UNII: 9BR6002P6E)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
WATER (UNII: 059QF0KO0R)	
CAPRYLYL GLUCOSIDE (UNII: V109WUT6RL)	
LAURETH-4 (UNII: 6HQ855798J)	
POLIHEXANIDE (UNII: 322U039GMF)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TRIMETHYL OCTADECYL AMMONIUM CHLORIDE (UNII: CZ70647U92)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51706-907-01	650 g in 1 TUBE; Type 0: Not a Combination Product	05/27/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/27/2020	

Labeler - Landy International (545291775)

Establishment

Name	Address	ID/FEI	Business Operations
Zhangzhou Kingbaby Health products Co.,Ltd		415797593	manufacture(51706-907)

Revised: 5/2020 Landy International