

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.

51706-906 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 irritation 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 Barbados leaf extract, Aqua, Citric Acid, Caprylyl Glucoside, Laureth-4, Polyhexanide, Phenoxyethanol, Triethoxysilylpropyl Steardimonium Chloride.

Package Label - Principal Display Panel



KILLS 99.99% OF HARMFUL GERMS





WITH SOOTHING ALOE VERA

SANITIZING HAND WIPES

LONG-LASTING, ALCOHOL-FREE PROTECTION FROM GERMS



FAST ACTING 15 SECOND FORMULA



80 WIPES

BLEACH FREE



100% LINT FREE

CAUTION: KEEP OUT OF THE REACH OF CHILDREN
SEE BACK PANEL FOR PRECAUTIONARY STATEMENTS AND FIRST AID



SANITIZING HAND WIPES

Please dispose of used wipes responsibly. Do not dispose used wipes down the toilet.

Drug Facts

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QUESTIONS? +1(888) 331-8332, M-F, 9:00AM-5:00PM (EST)



DISTRIBUTED BY: ESC BRANDS, LLC
664 OLD HARGRAVE ROAD, SUITE B
LEWINGTON, NC 27295
919-331-8332
www.my-shield.com

FORMULATED IN THE USA
MADE IN CHINA

E-1003-06



NET CONTENTS: MINIMUM WT. 1 lb. 4.3 oz. (0.65kg) **80 WIPES** 5.91 in x 7.09 in (15.0 cm x 18.0 cm)

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SANITIZING HAND WIPES				
benzalkonium chloride cloth				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51706-906	
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)				
POLYHEXANIDE (UNII: 322U039GMF)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
TRIMETHYLOCTADECYLAMMONIUM CHLORIDE (UNII: CZ70647U92)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51706-906-01	650 g in 1 TUBE; Type 0: Not a Combination Product	05/27/2020	
2	NDC:51706-906-02	120 g in 1 BAG; 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	
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Labeler - Landy International (545291775)

Establishment

Name	Address	ID/FEI	Business Operations
Zhangzhou Zhongnan Nursing Products Co.,ltd		416376835	manufacture(51706-906)

Revised: 5/2020

Landy International