BZK SANITIZING HAND WIPES- benzalkonium chloride cloth Premium PPE, LLC

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.

BZK Sanitizing Hand Wipes

Directions for Use:

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

Active Ingredient:

Benzalkonium Chloride 0.13%

Purpose:

Antiseptic/ hand and skin sanitizer

Uses:

Sanitizing hand wipes help decrease bacteria on the skin. Recommended for single use

Inactive Ingredients:

Aloe Barbadensis leaf extract, Aqua, Citric Acid, Caprylyl Glucoside, L a u r e t h - 4, P o l y h e x a n i d e , Phenoxyethanol, Trielhoxysilylpropyl Steardimonium Chloride.

Warnings:

Do not freeze

+ For external use only

Do not use

+ in ears, eyes or mouth

When using this product,

- +avoid contact with the eyes
- + In case of contact, flush eyes with water

Stop use and ask a doctor if

+redness or irritation develops and persists for more than 72 hours

Keep out of reach of children

+Children should be supervised when using this product.

Questions?

1-800-920-7650 Mon-Fri 10AM-4PM (EST)

Package Labeling:

BZK™

Sanitizing Hand Wipes

UP TO 4 HOURS OF PROTECTION + Kills up to 99.9% of Harmful Germs and Bacteria + Naturally Derived, Alcohol and Bleach Free + Gentle and Non-stinging with Aloe Vera

MADE IN THE USA

MIN WT 1 LB 4.3 OZ

80 WIPES 6 IN X 8 IN (16 CM X 20 CM)



B7.K™

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

Please dispose of used wipes responsibly. Do not dispose used wipes into toilets.



MADE IN THE USA

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BZK™ products are manufactured under strict quality guidelines under 21 CFR at a cGMP FDA-registered site.

Distributed By BZK Health Buffalo NY, 14221

bzkhealth.com



BZK SANITIZING HAND WIPES

benzalkonium chloride cloth

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:81529-003

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZ ALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
CAPRYLYL GLUCOSIDE (UNII: V109WUT6RL)	
LAURETH-4 (UNII: 6HO855798I)	

POLIHEXANIDE (UNII: 322U039GMF)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TRIETHOXYSILYLPROPYL STEARDIMONIUM CHLORIDE (LINII: XGN40Y0C7B)	

P	Packaging					
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:81529- 003-01	80 in 1 CANISTER	02/03/2020			
1		6.25 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)				

Marketing Information				
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
part333E	02/03/2020			
	Application Number or Monograph Citation	Application Number or Monograph Marketing Start Citation Date		

Labeler - Premium PPE, LLC (117835683)

Revised: 2/2021 Premium PPE, LLC