BZK SANITIZING HAND WIPE- 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 Wipe

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:

Cleans and sanitizes the skin and decreases bacteria. 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

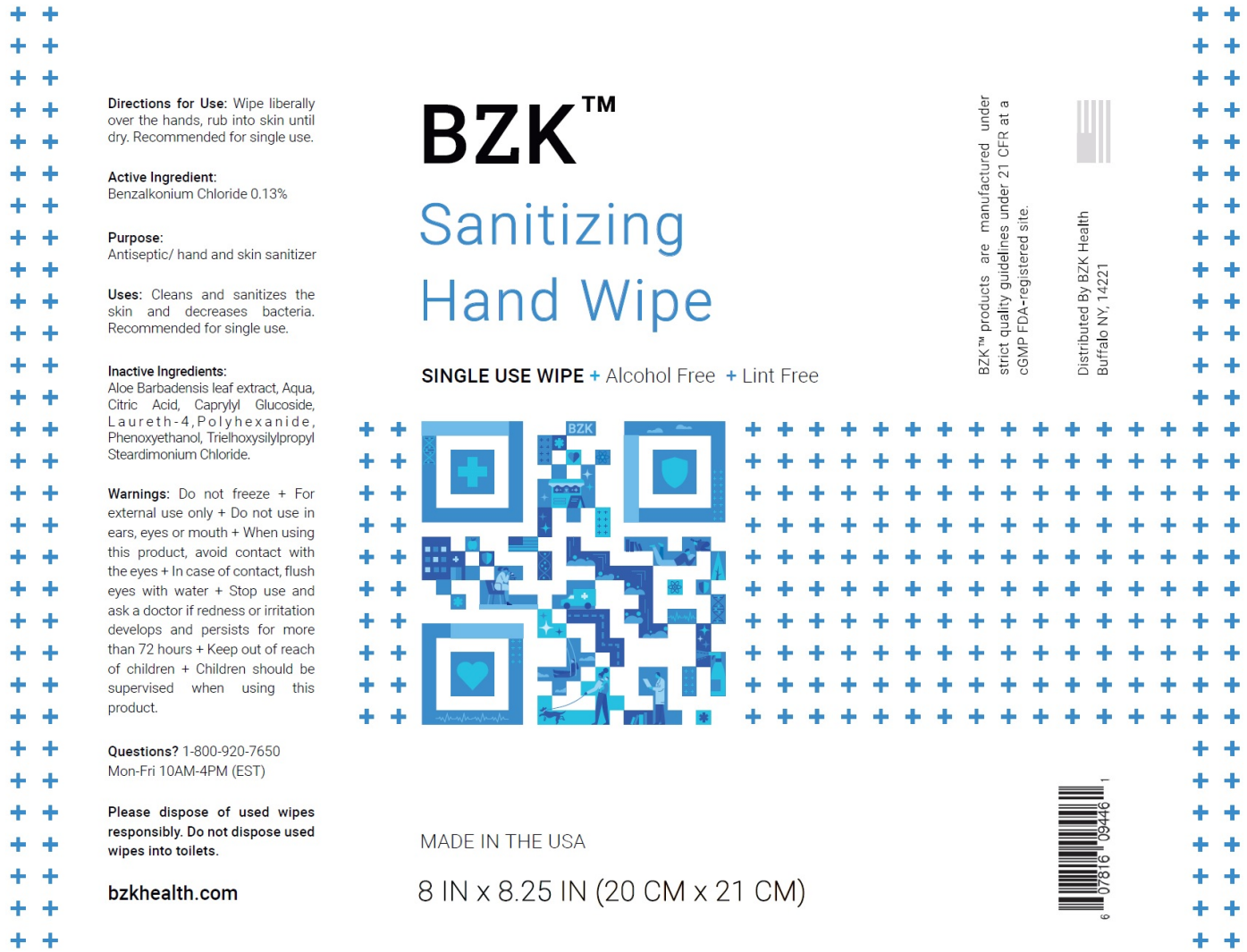
Keep out of reach of children

+Children should be supervised when using this product.

Questions?

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

Package Labeling:



BZK SANITIZING HAND WIPE

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81529-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM 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: 6HQ855798J)	
POLIHEXANIDE (UNII: 322U039GMF)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
TRIETHOXYSILYLPROPYL STEARDIMONIUM CHLORIDE (UNII: XGN40YQC7B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81529-004-01	1 in 1 PACKET	02/03/2021	
1		6.7 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	02/03/2021	

Labeler - Premium PPE, LLC (117835683)

Revised: 2/2021 Premium PPE, LLC