BZK ALCOHOL FREE HAND SANITIZER- benzalkonium chloride spray
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 Alcohol Free Hand Sanitizer

Drug Facts

Active Ingredient:

Benzalkonium Chloride 0.13%

Purpose:

Hand & Skin Sanitizer

Uses:

BZK™ Alcohol Free Hand Sanitizer provides revolutionary protection in an alcohol-free formula. The scientifically-proven solution kills harmful germs, bacteria, and microbes, and gently soothes and softens skin with aloe vera. Recommended for repeated use.

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.

Directions:

Apply liberally to the palms of the hands or areas of damaged skin. Rub into skin until dry. Recommended for repeated use.

Other Information:

Store in a cool dry place below 104°F(40°C).

Inactive Ingredients:

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

Questions?

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

Package Labeling:

BZK ALCOHOL FREE HAND SANITIZER

benzalkonium chloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81529-001
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:81529-001-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/03/2021	

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