BZK ALCOHOL FREE HAND SANITIZER- benzalkonium chloride liquid Sarati International, Inc.

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)

Product Label

BZK™

Alcohol Free Hand Sanitizer

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

1.7 FL OZ (50 ML)

MADE IN THE USA

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

1.7 FL OZ

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2 FL OZ

BZK™

Alcohol Free Hand Sanitizer

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MADE IN THE USA

2 FL OZ (59 ML)



Active Ingredient: Benzalkonium Chloride 0.13%

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BZK ALCOHOL FREE HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:67676-006

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)

BENZALKONIUM O.13 g in 100 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: XGN40YOC7B)				

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:67676- 006-01	50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/31/2021	
2	NDC:67676- 006-02	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/31/2021	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	505G(a)(3)	03/31/2021				

Labeler - Sarati International, Inc. (160219770)

Registrant - Sarati International, Inc. (160219770)

Establishment						
Name	Address	ID/FEI	Business Operations			
Sarati International, Inc.		160219770	pack(67676-006)			

Revised: 10/2023 Sarati International, Inc.