

HAND SANITIZING WIPES- benzalkonium chloride cloth
Ez Products Of South Florida, LLC

Hand Sanitizing Wipes

Drug Facts

Active ingredient

Benzalkonium Chloride (BZK) 0.1%

Purpose

Antibacterial

Use

- Decrease bacteria on the skin.

Warnings

For external use only.

When using this product

avoid contact with eyes.If contact occurs, rinse thoroughly with water.

Do not use

if irritation and redness develop.

Stop use and seek medical attention

if condition persists for more than 72 hours.

Keep out of reach of children.

If swallowed, seek medical attention or contact Poison Control Center.

DIRECTIONS

1. Locate tear-notch on the side of pouch. 2. Tear straight across to open. 3. Pull first wipe from center of roll and thread wipe through dispensing nozzle

Do not remove wipes roll from pouch.

Other information

- Dispose of wipe in the proper container.

- Do not flush down the toilet.

Inactive ingredients

2-Bromo-2-Nitropropane-1,3-Diol, Alcohol, Aloe Barbadensis Leaf Extract, Chamomilla Recutita Flower Extract, Citric Acid, Fragrance, Iodopropynyl Butylcarbamate, Lauryl Glucoside, Phenoxyethanol, Propylene Glycol, Tetrasodium EDTA, Water.

Package Labeling:



- KILLS 99.99% OF MOST COMMON GERMS
- PARABEN FREE

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HAND SANITIZING WIPES

1500 Wet Wipes • 6.3 in. x 6.7 in. (16 cm. x 17 cm.)

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Distributed by:
EZP International
Boca Raton, FL 33432
888-465-1001 • 561-362-2020
www.ezpint.com
Made in China



HAND SANITIZING WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69446-220
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM	1 mg

UNII:7N6JUD5X6Y)	CHLORIDE	in 1 g
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Inactive Ingredients

Ingredient Name	Strength
BRNOPOL (UNII: 6PU1E16C9W)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CHAMOMILE (UNII: FGL3685T2X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE SODIUM (UNII: MP1J8420LU)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69446-220-00	1500 in 1 BAG	04/30/2021	
1		2.346 g 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 Drug	505G(a)(3)	04/30/2021	

Labeler - Ez Products Of South Florida, LLC (113456060)

Revised: 10/2023

Ez Products Of South Florida, LLC