

ANTISEPTIC TOWELETTE- benzalkonium chloride cloth
Taizhou Kangping Medical Science And Technology Co., Ltd.

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.

Ever Ready
BZK Antiseptic Towelette

Active ingredients

Benzalkonium chloride, 0.13%

Purpose

Antiseptic

Uses

For handwashing to decrease bacteria on the skin.

Warnings

For external use only.

Do not use

in the eyes or apply over large areas of the body.

Stop use

if irritation, redness or other symptoms develop.

Consult a doctor if the condition persists or gets worse.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Open packet, unfold and use as a washcloth.

Inactive ingredients

Purified water

Other information

Store at room temperature: 15°-30°C (59°-86°F)

PROFESSIONAL USE

BZK Antiseptic Towelette

- Apply topically as needed
- Sterile unless pouch is opened or damaged

For External Use Only

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ER[®]

DISTRIBUTED BY:
EVER READY FIRST AID AND MEDICAL SUPPLY CO., LTD
300 LIBERTY AVE, BROOKLYN, NY 11207

Reorder No. 0100034

Drug Facts
Active ingredients Purpose Benzalkonium Chloride, 0.13%..... Antiseptic
Uses For handwashing to decrease bacteria on the skin.
Warnings For external use only.
Do not use in the eyes or apply over large areas of the body.
Stop use if irritation, redness or other symptoms develop. Consult a doctor if the condition persists or gets worse.
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LOT 220583459 2022-11 2025-10 Made in China

ANTISEPTIC TOWELETTE

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71310-004
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		
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71310-004-01	1.6 mL in 1 POUCH; Type 0: Not a Combination Product	10/28/2022	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	10/28/2022		

Labeler - Taizhou Kangping Medical Science And Technology Co., Ltd. (543429840)

Revised: 10/2022

Taizhou Kangping Medical Science And Technology Co., Ltd.