

VLITEPRO BZK ANTISEPTIC TOWELETTE- benzalkonium chloride cloth
Yiwu Ori-Power Medtech Co.,Ltd.

VLITEPRO BZK Antiseptic Towelette

Drug Facts

Active ingredient

Benzalkonium Chloride, 0.13%

Purpose

Antiseptic

Use

First Aid to help reduce the risk of infection in minor cuts, scrapes and burns.

Warnings

For external use only.

Do not use

in the eyes or over large areas of the body

Ask a doctor before use if you have

deep or puncture wounds or serious burns.

Stop use and ask a doctor if

- 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

Clean the affected area

Inactive ingredients

Sodium Bicarbonate, Purified Water

VLITEPRO BZK ANTISEPTIC TOWELETTE

VLITEPRO[®]

BZK ANTISEPTIC
TOWELETTE

1
pad

DRUG FACTS

ACTIVE INGREDIENT

Benzalkonium Chloride 0.13%

PURPOSE

First Aid Antiseptic

USES

For handwashing to decrease bacteria on the skin

WARNINGS

For external use only

KEEP OUT OF REACH OF CHILDREN

Made in China

NDC:

DRUG FACTS (CONTINUED)

DO NOT USE

In the eyes or over large areas of the body

STOP USE

If irritation and redness develop, or if other symptoms occur, consult a physician immediately

DIRECTIONS

Tear open packet and use towelette to cleanse skin area.

Discard towelette appropriately after single use

OTHER INFORMATION

Store at room temperature

INACTIVE INGREDIENTS

Purified Water, Sodium Bicarbonate

Manufactured by:

Changzhou Maokang Medical Products Co., Ltd

Add: No. 88 of Longxi Avenue, Zhulin Town, Jintan District, Changzhou City, Jiangsu Province, China P. C.: 213200

LOT:

EXP:

VLITEPRO BZK ANTISEPTIC TOWELETTE

benzalkonium chloride cloth

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72459-020	
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)				
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72459-020-01	1 in 1 PACKAGE	05/19/2025	
1		1 in 1 POUCH		
1		1.6 mL in 1 POUCH; Type 0: Not a Combination		

1	Product		
Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	05/19/2025	

Labeler - Yiwu Ori-Power Medtech Co.,Ltd. (560451976)

Revised: 5/2025

Yiwu Ori-Power Medtech Co.,Ltd.