

**MEDICHOICE BZK ANTISEPTIC- benzalkonium chloride cloth
O&M HALYARD, INC.**

MediChoice BZK Antiseptic Towelette

Drug Facts

Active ingredient

Benzalkonium Chloride, 0.13% w/w

Purpose

First Aid Antiseptic

Use

First aid antiseptic product to apply topically to the skin to help prevent infection in minor cuts, scrapes, and burns.

Warnings

For external use only. Do not use on mucous membranes or broken or irritated skin. Do not use

Do not use

- in the eyes or apply over large areas of the body.
- longer than 1 week unless directed by a doctor.
- no infants. **Consult a doctor**
- in case of deep or puncture wounds, animal bites, or serious burns.

Stop use and 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

- Clean the affected area 1 to 3 times daily or as directed by a doctor. • May be covered with a sterile bandage when dry.

Other information

- Store at room temperature.
- Allow pad to air dry before disposal.

Inactive ingredient

Purified Water, Sodium Bicarbonate

Package Labeling:

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NDC 39892-0304-1

MEDICHOICE[®]

BZK Antiseptic Towelette

Benzalkonium Chloride First Aid Antiseptic

100 Each/Box

Reorder: **BZK100**



Single Use Only



DO NOT
RE-STERILIZE

NOT MADE WITH
natural rubber latex

sterile^R

Distributed by Owens & Minor
9120 Lockwood Boulevard
Mechanicsville, VA 23116
Made in China

WARNING: For external use only.
CAUTION: Sterility of contents guaranteed in
unopened, undamaged package.



(01) 3 0885632 10841 3

MEDICHOICE BZK ANTISEPTIC

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:39892-0304
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

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:39892-0304-1	10 in 1 CASE	08/29/2017	
1		100 in 1 BOX		
1		1 in 1 PACKET		
1		0.75 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	M003	08/29/2017	

Labeler - O&M HALYARD, INC. (081057389)**Registrant** - Owens & Minor Inc. (847412269)

Revised: 10/2025

O&M HALYARD, INC.