# EASY CARE BZK ANTISEPTIC TOWELETTE- benzalkonium chloride swab Adventure Ready Brands

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.

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#### **Easy Care BZK Antiseptic Towelette**

### **Drug Facts**

#### **Active Ingredient**

Benzalkonium Chloride, 0.133%

#### **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

**REF** 855

855 NDC 44224-0113-0



# BZK ANTISEPTIC TOWELETTE

Contains Benzalkonium Chloride



Manufactured for:



Made in China





944 Industrial Park Rd. Littleton, NH 03561 easycarefirstaid.com

5020-0113-D

# **Drug Facts**

**Active ingredient**Benzalkonium Chloride, 0.133% Antiseptic **Purpose** Use First Aid to help reduce the risk of infection in minor cuts, scrapes and burns.

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Poison Control Center right away

**Directions** Clean the affected area

**Inactive ingredients** Sodium Bicarbonate, Water

EXP:

LOT

### EASY CARE BZK ANTISEPTIC TOWELETTE

**TOPICAL** 

benzalkonium chloride swab

<b>Product</b>	Intorma	ation
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**Route of Administration** 

**HUMAN OTC DRUG** NDC:44224-0113 Item Code (Source) **Product Type** 

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZ ALKONIUM CHLORIDE	1.33 mg in 1 mL		

Inactive Ingredients			
Ingredient Name	Strength		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)			
WATER (UNII: 059QF0KO0R)			

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:44224- 0113-0	1.4 mL in 1 POUCH; Type 0: Not a Combination Product	01/03/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/03/2022	

# **Labeler -** Adventure Ready Brands (064437304)

# Registrant - Adventure Ready Brands (064437304)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Adventure Ready Brands		064437304	manufacture(44224-0113)	

Revised: 12/2021 Adventure Ready Brands