FIRST AID DIRECT ANTISEPTIC WIPE- benzalkonium chloride sponge
Cintas Corporation

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.

First Aid Direct Antiseptic Wipe

Active ingredient (in each towelette)

Benzalkonium Chloride 0.133%

Purpose

first aid antiseptic

Uses

first aid to reduce bacteria in minor cuts, scrapes and burns

Warnings

For external use only

Do not use
- in the eyes or apply over large areas of the body
- longer than 1 week unless directed by a doctor

Ask a doctor before use if you have
- deep or puncture wounds
- animal bites
- 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
- tear open packet, unfold, use and discard
- clean the affected area
- maybe covered with a sterile bandage. If bandaged, let dry first.

Other information
- do not use if packet is torn, cut or opened
- store at room temperature

Inactive ingredients

purified water, sodium bicarbonate

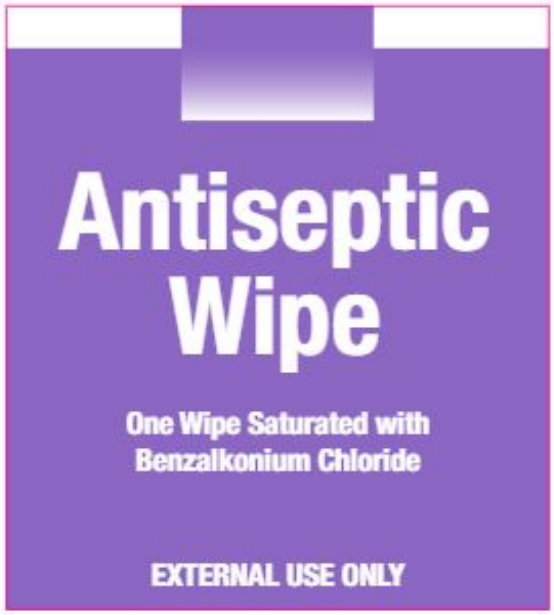
Questions? 1-877-973-2811

Principal Display Panel

Antiseptic Wipe

One Wipe Saturated with Benzalkonium Chloride

EXTERNAL USE ONLY



FIRST AID DIRECT ANTISEPTIC WIPE

benzalkonium chloride sponge

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42961-127
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	.133 mg in 100 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
	NDC:42961-127			

1	NDC:42961-127-02	10 in 1 BAG	09/28/2023	
1	NDC:42961-127-01	1.4 mL in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:42961-127-03	100 in 1 BOX	09/28/2023	
2	NDC:42961-127-01	1.4 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/28/2023	

Labeler - Cintas Corporation (056481716)

Establishment

Name	Address	ID/FEI	Business Operations
Changzhou Maokang Medical Products Co., Ltd.		421317073	manufacture(42961-127)

Revised: 9/2023 Cintas Corporation