ANTISEPTIC BANDAGES- benzalkonium chloride dressing
Pharmaplast SAE

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.

Cure Aid Topical antimicrobial adhesive bandages

Drug Facts

Active ingredients

Benzalkonium Chloride 0.1%

Purpose

Topical antimicrobial adhesive bandage

Uses

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

Warnings

- **For external use only.**
- **Stop use and consult a doctor if** the condition persists or gets worse. Do not use longer than 1 week unless directed by a doctor.
- **Keep out of reach of children.** If swallowed, get medical help or contact Poison Control Center right away.

Directions

Clean and dry the affected area. Apply a sterile bandage on the area 1 to 3 times daily.

Inactive Ingredients

None

CURE-AID®

INTENDED USE:
Antimicrobial Adhesive Bandages are to be applied to the skin for topical application. The bandages help provide an antibacterial barrier for minor cuts and scrapes.

DIRECTIONS:
Apply bandages to clean, dry skin. Change daily or when pad becomes wet. Sterile unless wrapper is damaged or open.

WARNINGS:
FOR MEDICAL EMERGENCIES SEEK PROFESSIONAL HELP

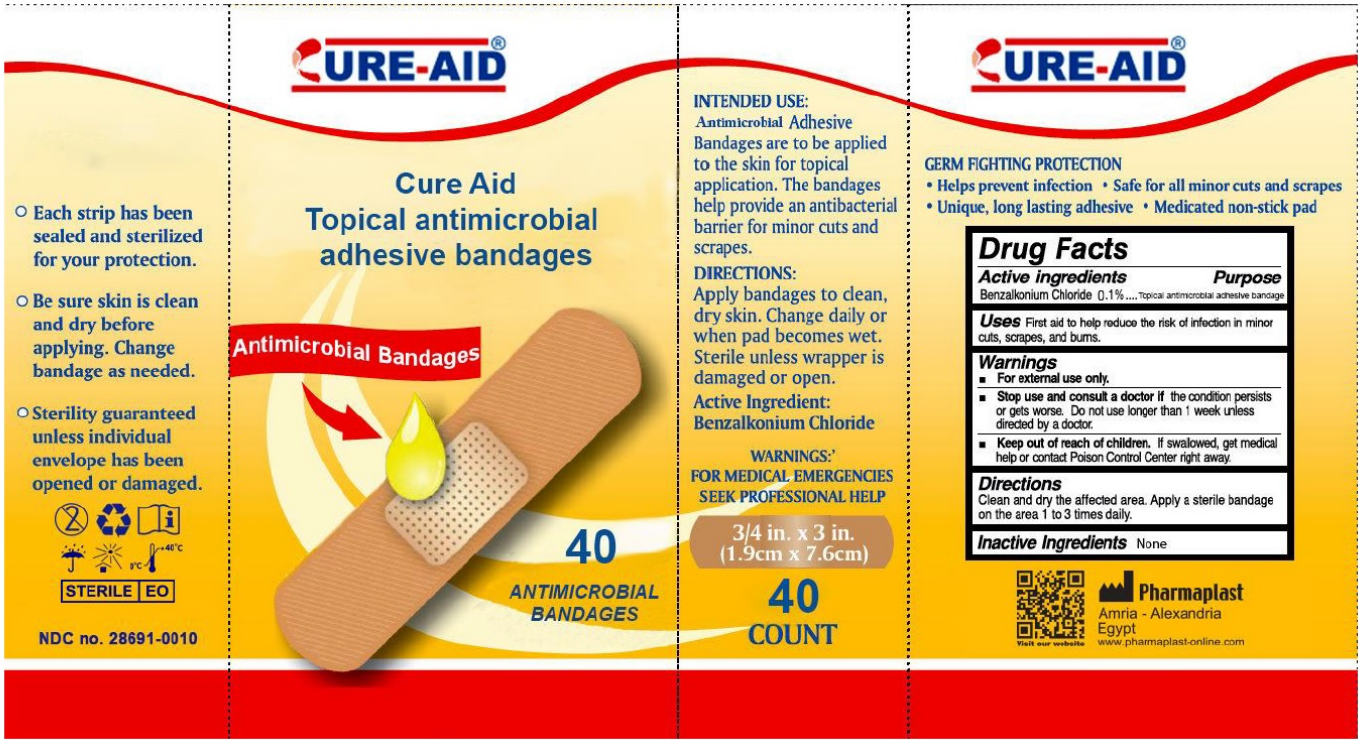
GERM FIGHTING PROTECTION
• Help prevent infection • Safe for all minor cuts and scrapes • Unique, long lasting adhesive • Medicated non-stick pad

- **Each strip has been sealed and sterilized for your protection.**
- **Be sure skin is clean and dry before applying. Change bandage as needed.**
- **Sterility guaranteed unless individual envelope has been opened or damaged.**

Pharmaplast

Amria – Alexandria
Egypt
www.Pharmaplast-online.com

Packaging



ANTISEPTIC BANDAGES

benzalkonium chloride dressing

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:28691-0010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 mg in 100 mg

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
Does not contain	**NATURAL LATEX RUBBER** (UNII: 2LQ0UUW8IN)	0 mg in 100 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:28691-0010-5	20 in 1 BOX	10/07/2016	
1		0.033 mg in 1 PATCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		
2	NDC:28691-0010-6	30 in 1 BOX	10/07/2016	
2		0.033 mg in 1 PATCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		
3	NDC:28691-0010-7	40 in 1 BOX	10/07/2016	
3		0.033 mg in 1 PATCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/07/2016	

Labeler - Pharmaplast SAE (644773319)

Establishment

Name	Address	ID/FEI	Business Operations
Pharmaplast SAE		644773319	analysis(28691-0010) , label(28691-0010) , manufacture(28691-0010) , pack(28691-0010)

Revised: 11/2020 Pharmaplast SAE