GREEN GUARD FIRST AID ANTISEPTIC- benzalkonium chloride spray Ultra Distributors Inc

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.

GREEN GUARD® Antiseptic spray

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

First aid antiseptic

Uses

First aid to help prevent infection in minor cuts, scrapes & burns

Warnings

For external use only.

Flammable, keep away from fire or flame.

Do not use

- near eyes or mucous membranes
- on deep or puncture wounds, animal bites, or serious burns
- over large areas of the body
- more than one week unless directed by a doctor

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 affected area & spray 1 to 3 times daily
- may be covered with a sterile bandage
- not to be used on children under 12 years of age

Inactive ingredients

ethyl alcohol, purified water

Questions or comments?

1-800-869-6970

Treats minor cuts, scrapes and abrasions

Helps prevent infection

Store at 68°-77°F (20°-25°C)



GREEN GUARD FIRST AID ANTISEPTIC

benzalkonium chloride spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78495-112
Route of Administration	TOPICAL		

	Active Ingredient/Active Moiety		
ı	Ingredient Name	Basis of Strength	Strength
	BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

ALCOHOL	(IINII)	3K99	58 V	790M)

ı	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:78495-112- 01	59.1 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/19/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/19/2020	

Labeler - Ultra Distributors Inc (007160073)

Revised: 12/2020 Ultra Distributors Inc