OMNICIDE ANTIMICROBIAL- benzalkonium chloride gel
SteriWeb Medical, LLC

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.

STERIWEB RX (as PLD) - OMNICIDE ANTIMICROBIAL GEL (69085-466-05)

Active Ingredient

Benzalkonium Chloride 0.13% w/w

Purpose

First Aid Antiseptic

Uses

First aid to help treat, protect and prevent skin infection associated with
- minor cuts
- scrapes
- burns

Warnings

For external use only

Ask a doctor before use if you have
- deep or puncture wounds
- animal bites
- serious burns

When using this product
- Do not use in or near the eyes
- Do not use if you have a history of allergy to any of the ingredients

Stop and ask a doctor if
- condition worsens
- symptoms persist for more than 7 days, or clear up and occur again within a few days.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away

Directions

- adults and children 2 years and older: clean the affected area; apply a small amount on the area 1-3 times daily; may be covered with a sterile bandage (let dry first)
- children under 2 years: ask a doctor

Other information

Avoid excessive heat.

Inactive Ingredients

USP Petrolatum, Water, Polihexanide (PHBM)

Questions?

1-360-448-7881

or www.steriwebrx.com

OMNICIDE ANTIMICROBIAL

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69085-466
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)	
POLIHEXANIDE (UNII: 322U039GMF)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69085-466-05	30 g in 1 TUBE; Type 0: Not a Combination Product	05/07/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/07/2016	

Labeler - SteriWeb Medical, LLC (079409860)

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