KENDALL ANTIMICROBIAL CLEANSER - benzalkonium chloride liquid
Covidien 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.

Kendall Antimicrobial Cleanser

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Use

- a no-rinse topical antiseptic that protects against microbial contamination
- ideal for chapped or lightly traumatized skin

Warnings
For external use only.

When using this product

- do not get into eyes

Do not use on

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if condition worsens or symptoms last more than seven days.

Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- spray soiled or odorous areas
- gently wipe clean. Repeat as necessary.
- pat dry. No rinse necessary.
- apply Kendall Moisturizing Lotion, Moisture Barrier Cream or Soothing Ointment as necessary

Other Information

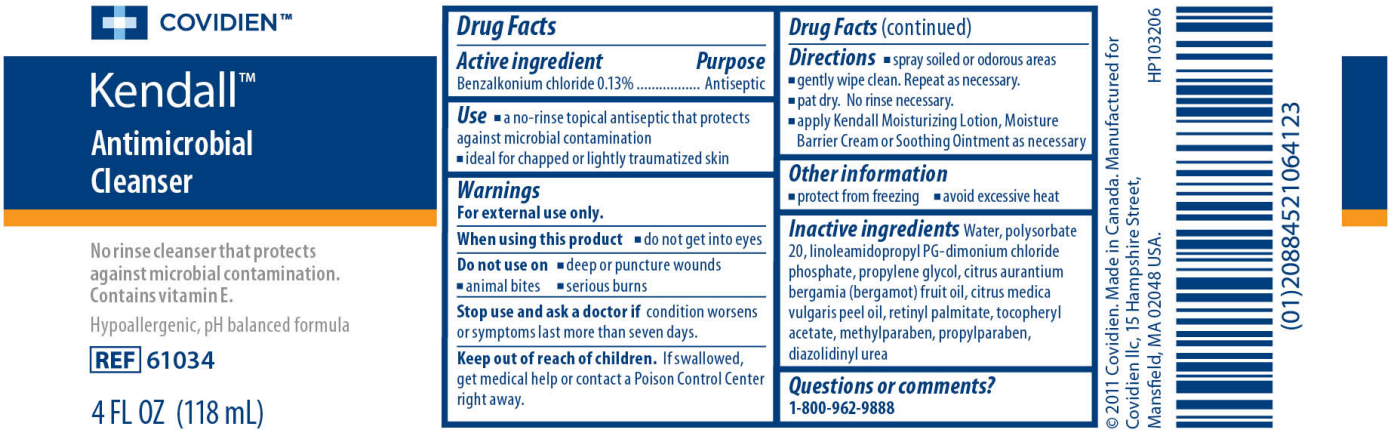
- protect from freezing
- avoid excessive heat

Inactive ingredients

Water, Polysorbate 20, linoleamidopropyl PG-dimonium chloride phosphate, propylene glycol, cirtus aurantium bergamia (bergamot) fruit oil, citrus medica vulgaris peel oil, retinyl palmitate, tocopheryl acetate, methylparaben, propylparaben, diazolidinyl urea

Questions or comments? 1-800-962-9888

Image of 4 Ounce Label



KENDALL ANTIMICROBIAL CLEANSER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:28851-691
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 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PHOSPHATE ION (UNII: NK08V8K8HR)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BERGAMOT OIL (UNII: 39W1PKE3JI)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
ACETATE ION (UNII: 569DQM74SC)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:28851-691-01	118 mL in 1 BOTTLE, SPRAY		
2	NDC:28851-691-02	236 mL in 1 BOTTLE, SPRAY		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/05/2011	

Labeler - Covidien Inc. (805770828)

Revised: 4/2011 Covidien Inc.