ROYAL NON-ALCOHOL FOAM HAND SANITIZER- benzalkonium chloride solution
CWGC LA Inc.

CWGC (as PLD) - Royal Non-Alcohol Foam Hand Sanitizer (70415-205)

Active ingredient

BENZALKONIUM CHLORIDE 0.13%

Purpose

Antibacterial

Uses

- For handwashing to reduce bacteria on the skin. Recommended for repeated use.

Warnings

For external use only

Avoid contact with eyes. In case of eye contact, flush eyes with water.

Keep out of reach of children.

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

Stop use and ask a doctor if irritation or redness develops and persists.

Directions

- Apply foam sanitizer to hands.
- Rub over surfaces of both hands for 15 seconds.
- No rinsing required.

Inactive ingredients

Water, Coco-Glucoside, Laurtrimonium Chloride, Cocamidopropylamine Oxide, Citric Acid, Fragrance.

ROYAL NON-ALCOHOL FOAM HAND SANITIZER

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70415-205
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 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	

TROLAMINE (UNII: 9O3K93S3TK)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70415-205-01	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/22/2017	

Labeler - CWGC LA Inc. (034967904)

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