

**SIERRA SOFT NON-ALCOHOL HAND SANITIZER FOAM- benzethonium chloride liquid
CWGC LA 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.

CWGC (as PLD) - Sierra Soft Non-Alcohol Hand Sanitizer Foam (70415-104)

Drug Facts

Active ingredient

BENZETHONIUM CHLORIDE 0.25%

Purpose

Antibacterial

Uses

- For sanitizing to reduce bacteria on the skin.

Warnings

For external use only

When using this product

- avoid contact with eyes. In case of eye contact, flush with water

Keep out of reach of children.

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

Directions

- Apply one shot of foaming sanitizer to dry hands
- Rub into skin
- No rinsing required

Inactive ingredients

WATER, PROPYLENE GLYCOL, COCAMIDOPROPYL BETAINE, COCAMINE OXIDE, DISODIUM EDTA, DMDM HYDANTOIN, TRIETHANOLAMINE.

Package Labeling:



FOAM

Non-Alcohol Hand Sanitizer

Drug Facts

Active ingredient	Purpose
Benzethonium Chloride 0.25%	Antibacterial

Use ■ For sanitizing to reduce bacteria on the skin

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Directions

- Apply one shot of foaming sanitizer to dry hands
- Rub into skin
- No rinsing required

Inactive ingredients

Water, Propylene Glycol, Cocamidopropyl Betaine, Cocamine Oxide, Disodium EDTA, DMDM Hydantoin, Triethanolamine.

1L 2
33.8 fl. oz.



Manufactured for
Sierra Health Corp.
Placentia, CA 92870
855-686-1888
Made in PRC

SIERRA SOFT NON-ALCOHOL HAND SANITIZER FOAM

benzethonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70415-104
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	2.5 g in 1 L
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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)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70415-104-11	1 L in 1 BOTTLE; Type 0: Not a Combination Product	12/13/2016	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/13/2016	

Labeler - CWGC LA Inc. (034967904)