BROMPTON AND LANGLEY MOISTURIZING HAND SANITIZER MARKET STRAWBERRY- benzalkonium chloride gel Cita International Limited

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.

Active Ingredient

Alcohol 62%

□**Purpose:**□ Antiseptic

 \Box Use

To decrease bacteria on the skin that could cause disease.

Keep out of reach of children.

Warnings

- Keep away from flame or fire
- for external use only-hands.
- keep out of eyes.
- stop use and ask a Doctor if irritation or redness persists for more than 72 hours
- do not inhale or ingest, if swallowed, get medical help or contact a poison control center right away.

Directions

- spray a thumb size amount into hands.
- rub hands together gently making sure to cover all areas in between fingers
- let hands air dry

Inactive Ingredients

water (aqua/eau), glycerin, propylen glycol, Isopropyl alcohol, phenoxythenol, acrylates,

lauryl glycol ether, carbomer, Vitamin E, fragrance (parfum).

Brompton and Langley Strawberry Scentend Hand Sanitizer 15mL 7/11/2016 image022.png



STRAWBERRY

benzalkonium chloride gel

Product Informati	on
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Product Type HUMAN OTC DRUG NDC:70805-001 Item Code (Source)

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.26 g in 26 g

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Inactive Ingredients

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Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6 A3C0 OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PHENO XYETHANOL (UNII: HIE49 2ZZ3T)	
METHO XY PEG-40 (UNII: 6 AXS45P1QU)	
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)	

Product Characteristics

.ALPHA.-TO COPHEROL ACETATE (UNII: 9E8X80D2L0)

1 Totalet Characteristics			
Color		Score	
Shape		Size	
Flavor	STRAWBERRY	Imprint Code	
Contains			

Packaging

	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:70805-001- 01	15 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/11/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/03/2016	

Labeler - Cita International Limited (665926739)

Establishment				
Name	Address	ID/FEI	Business Operations	
Cita International Limited		665926739	manufacture(70805-001)	

Revised: 7/2016 Cita International Limited