

**BROMPTON AND LANGLEY COCONUT WATER AND HABISCUS HAND
SANITIZER COCONUT WATER AND HABISCUS- alcohol denat 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

☐**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

Coconut Water and Habiscus Scentend Hand Sanitizer

15mL



SANITIZER COCONUT WATER AND HABISCUS

alcohol denat gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70805-003
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

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
METHOXY PEG-40 (UNII: 6AXS45P1QU)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	COCONUT (Coconut Water and Habiscus)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70805-003-01	15 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/12/2016	

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(70805-003)

Revised: 7/2016

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