BROMPTON AND LANGLEY VANILLA MARSHMALLOW HAND SANITIZER VANILLA MARSHMALLOW- 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
Vanilla Marshmallow Scentend Hand Sanitizer
15mL

BROMPTON AND LANGLEY VANILLA MARSHMALLOW HAND SANITIZER

VANILLA MARSHMALLOW

alcohol denat gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70805-007
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	MARSHMALLOW (Vanilla Marshmallow)	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70805-007-01	15 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/13/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/13/2016	

Labeler - Cita International Limited (665926739)

Establishment

Name	Address	ID/FEI	Business Operations
Cita International Limited		665926739	manufacture(70805-007)