

**CABANA BAY BEACH RESORT HAND SANITIZER GEL- ethyl alcohol gel**  
**Carretta USA, 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.*

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**Universal's Cabana Bay Beach Resort Hand Sanitizer Gel**

**Drug Facts**

**Active Ingredient**

Ethyl Alcohol 75% v/v

**Purpose**

Antiseptic

**Use[s]**

Hand sanitizer to help reduce bacteria that potentially can cause disease.

For use when soap and water are not available.

**Warnings**

**For external use only. Flammable.**

**Keep away from heat or flame.**

**Do not use**

- in children less than 2 months of age
- on open skin wounds

**When using this product** keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

**Stop use and ask a doctor** if irritation or rash occurs. These may be signs of a serious condition.

**Keep out of reach of children.**

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

**Directions**

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

**Other information**

- Store between 15-30C (59-83F)
- Avoid freezing and excessive heat above 40C (104F)

**Inactive Ingredients:**

Water,Glycerin,Propylene glycol,Carbomer,Aloe Vera Extract,Triethanolamine,Disodium Edta

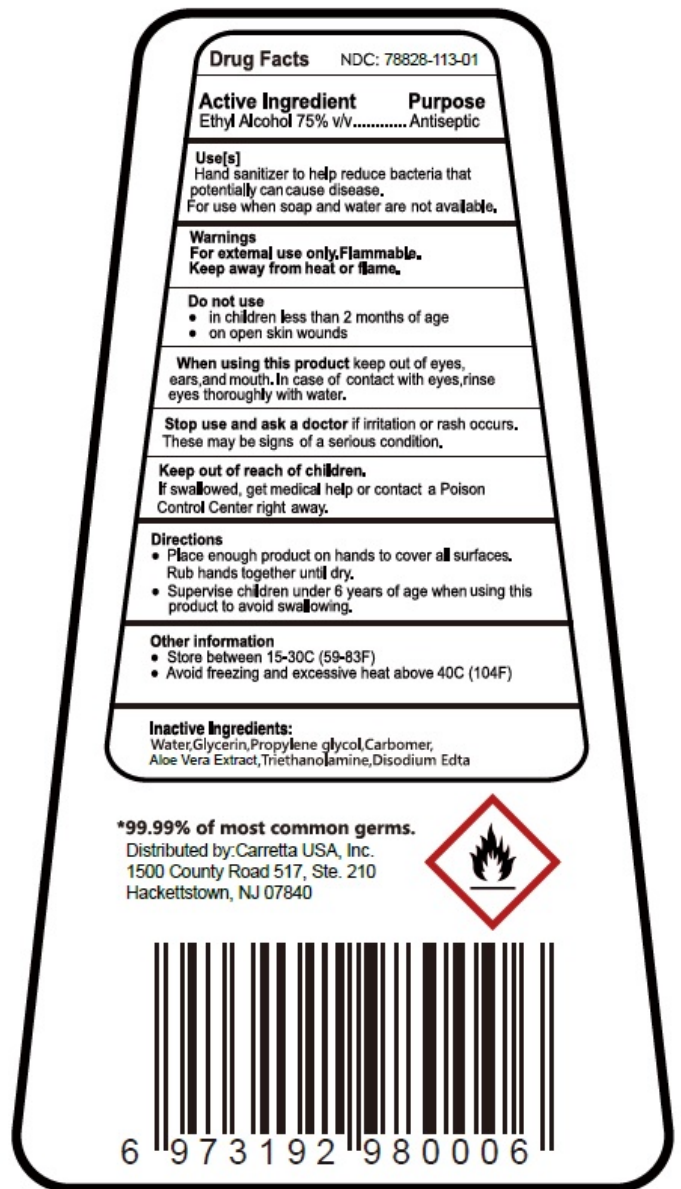
**\*99.99% of most common germs.**

Distributed by:Carretta USA, Inc.

1500 County Road 517, Ste. 210

Hackettstown, NJ 07840

**Packaging**



**CABANA BAY BEACH RESORT HAND SANITIZER GEL**

ethyl alcohol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:78828-113
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.75 mL in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
EDETATE DISODIUM ANHYDRO US (UNII: 8NLQ36F6MM)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78828-113-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/12/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/12/2020	

**Labeler** - Carretta USA, Inc (010851173)

Revised: 6/2020

Carretta USA, Inc