## LOEWS PORTOFINO BAY 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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## Loews Portofino Bay Hand Sanitizer Gel

## **Drug Facts**

## **Active Ingredient**

Ethyl Alcohol 75% v/v

## **Purpose**

Antiseptic

#### Uses[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)

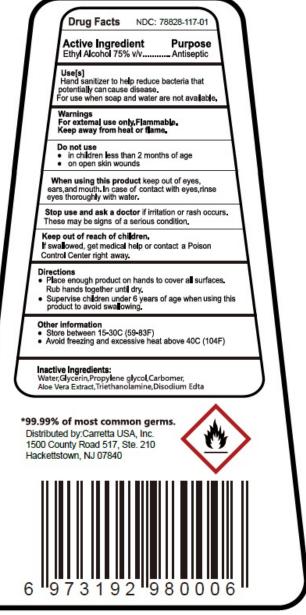
### **Inactive Ingredients:**

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

Distributed by:Carretta USA, Inc. 1500 County Road 517, Ste. 210 Hackettstown, NJ 07840

## **Packaging**





# LOEWS PORTOFINO BAY HAND SANITIZER GEL

ethyl alcohol gel

## **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:78828-117

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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 HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
TROLAMINE (UNII: 9O3K93S3TK)		
EDETATE DISO DIUM ANHYDRO US (UNII: 8 NLQ36 F6 MM)		

ı	Packaging					
ı	# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>		
ı	1 NDC:78828-117-	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