**SANIMAX HAND SANITIZER- alcohol liquid**
**Sanimax LLC**

*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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**Sanimax Hand Sanitizer**

**Drug Facts**

***Active ingredient[s]***

Alcohol 70% 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 usin this product**

keep out of eyes, ears, and mouth. In case de of contact with eyes, rinse eyes thoroughly with water.

**Stop use and ask a doctor**

if irritation or rash occurs. Theses may be signs of a serious condition.

**Keep out of reach of children.**

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

***Directions***

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

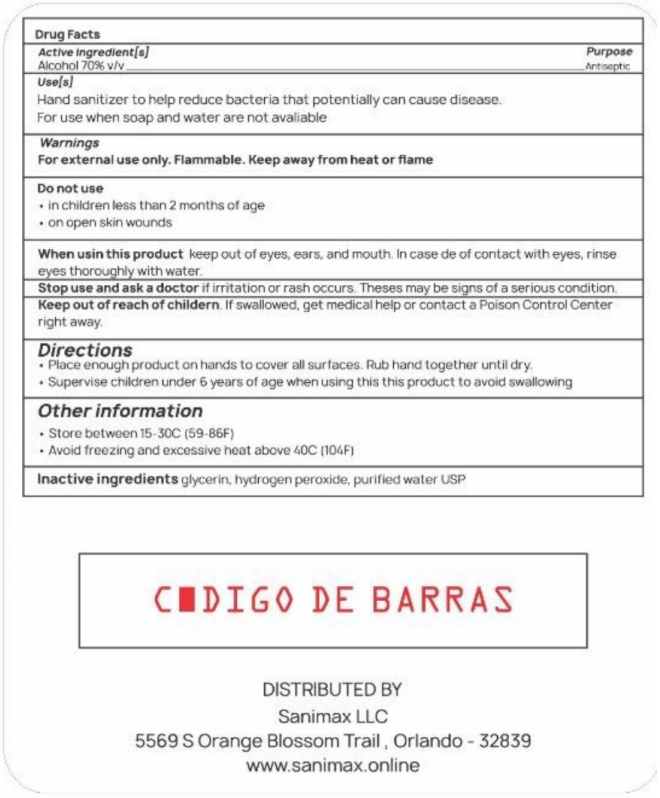
***Other information***

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

**Inactive ingredients**

glycerin, hydrogen peroxide, purified water USP

**Package Labeling:**


## SANIMAX HAND SANITIZER

alcohol liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:80289-001 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **ALCOHOL** (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 0.7 mL in 1 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **HYDROGEN PEROXIDE** (UNII: BBX060AN9V) | |
| **WATER** (UNII: 059QF0KO0R) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |

| | | | | |
|---|---|---|---|---|
| **1** | NDC:80289-001-00 | 500 mL in 1 BOTTLE; Type 0: Not a Combination Product | 09/03/2020 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333E | 09/03/2020 | |

## Labeler - Sanimax LLC (117576270)

Revised: 9/2020 Sanimax LLC