**HAND SANITIZER- alcohol liquid**
**Elevate Skincare, 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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**Elevate Hand Sanitizer**

***Active Ingredient***

Alcohol 62%

***Purpose***

Antiseptic

***Uses***

■ hand sanitizer to help reduce bacteria on the skin

***Warnings***

**Flammable. Keep away from fire or flame. For external use only**

**When using this product**

■ avoid getting into the eyes. ■ In case of eye contact immediately flush eyes thoroughly with water

**Stop use and ask a doctor if**

■ if irritation or redness develops ■ condition persists for more than 72 hours ■ redness is present

***Keep out of reach of children***

In case of accidental ingestion, contact a doctor or Poison Control Center immediately.

***Directions***

■ place enough product in your palm to thoroughly cover both hands ■ rub hands together until dry ■ no rinsing required

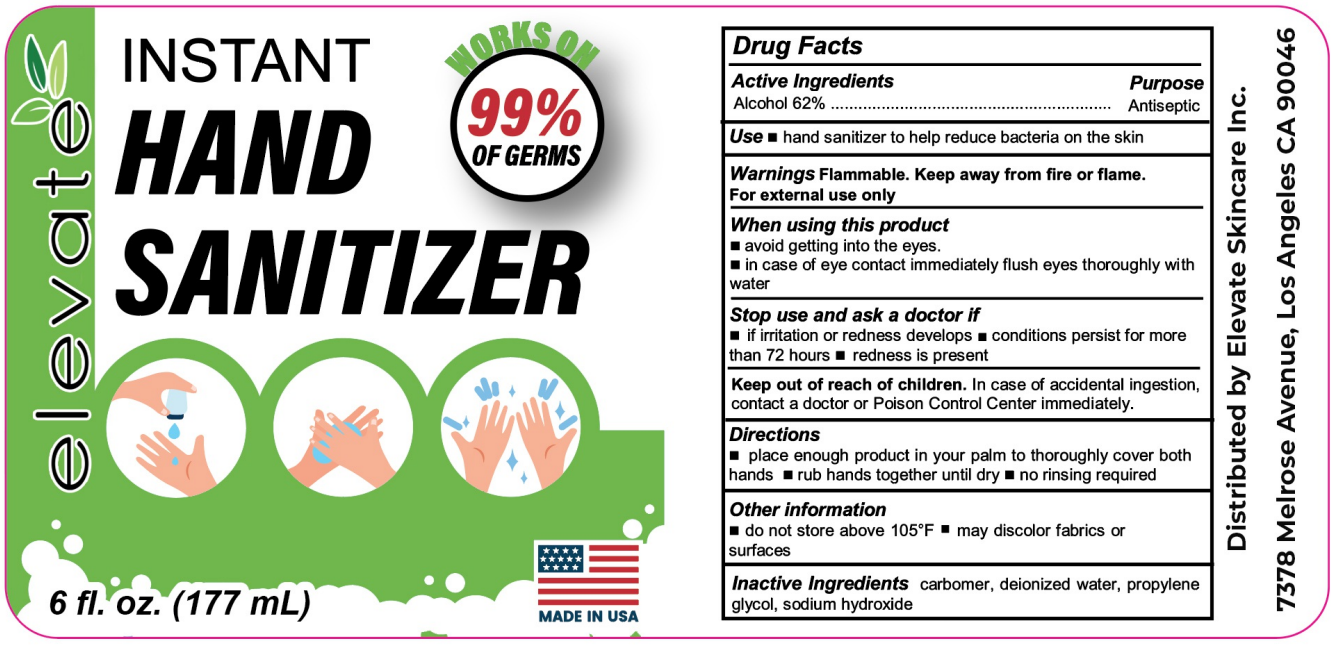
***Other information***

■ do not store above 105ºF ■ may discolor some fabrics or surfaces

***Inactive ingredients***

carbomer, deionized water, propylene glycol, sodium hydroxide

**Package Label**




# HAND SANITIZER

alcohol liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:74175-001 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **ALCOHOL** (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 62 mL in 100 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE** (UNII: 0A5MM307FC) | |
| **SODIUM HYDROXIDE** (UNII: 55X04QC32I) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:74175-001-06 | 177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 04/08/2020 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part333A | 04/08/2020 | |

## Labeler - Elevate Skincare, Inc. (111407277)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| IBG Holdings Inc. | | 138884643 | manufacture(74175-001) |

Revised: 4/2020 Elevate Skincare, Inc.