# SHIMMY ORIGINAL HAND SANITIZER- alcohol spray Shimmy Products 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.

-----

#### SHIMMY ORIGINAL HAND SANITIZER

### **Active Ingredients**

Ethyl Alcohol 70% v/v

#### **Purpose**

**Antimicrobial** 

#### Uses

- hand sanitizer to help reduce bacteria on the skin that could 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

- on 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 appears or lasts. These 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**

- Put enough product in your palm to cover your hands and rub hands together briskly until dry
- Children under 6 years of age should be supervised when using this product.

#### Other information

- Avoid freezing and excessive heat above 40°C (104°F).
- May discolor certain fabrics, materials or surfaces.

### **Inactive ingredients**

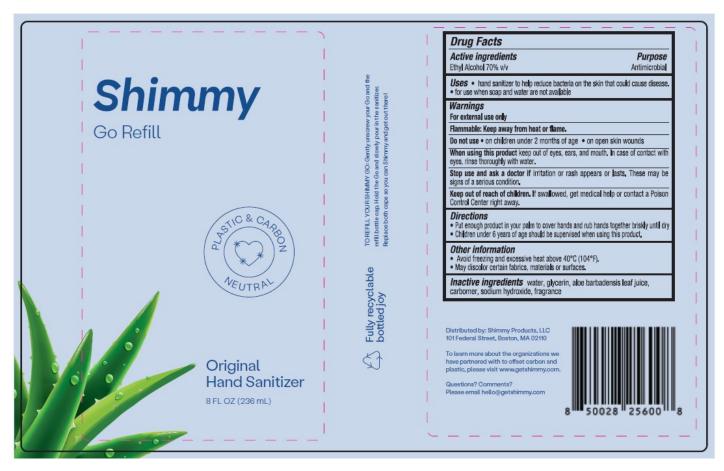
water, glycerin, aloe barbadensis leaf juice, carbomer, sodium hydroxide, fragrance

### Package Label - Principal Display Panel

SHIMMY ORIGINAL HAND SANITIZER

8 fl oz (236 mL)

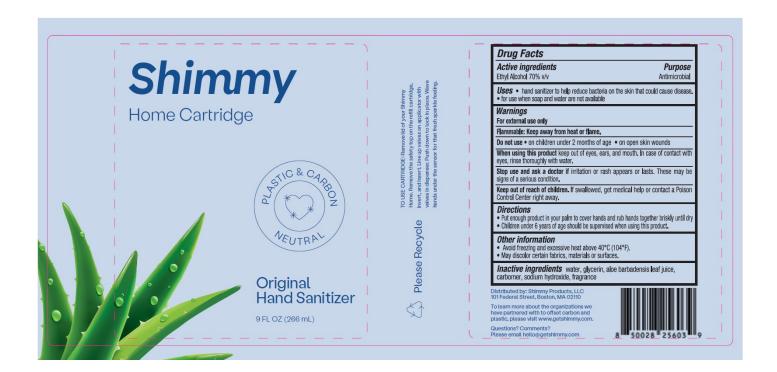
NDC: 81755-001-92



SHIMMY ORIGINAL HAND SANITIZER

9 fl oz (266 mL)

NDC: 81755-001-93



### SHIMMY ORIGINAL HAND SANITIZER

alcohol spray

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:81755-001

Route of Administration TOPICAL

### **Active Ingredient/Active Moiety**

Ingredient Name

Basis of Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

ALCOHOL (UNII: 3K9958V90M) ALCOHOL 70 mg in 100 mL

Inactive Ingredients				
Ingredient Name	Strength			
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
GLYCERIN (UNII: PDC6A3C0OX)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
WATER (UNII: 059QF0KO0R)				
CARBOMER 940 (UNII: 4Q93RCW27E)				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:81755-001- 92	236 mL in 1 CAN; Type 0: Not a Combination Product	06/14/2021		
2	NDC:81755-001- 93	266 mL in 1 CAN; Type 0: Not a Combination Product	06/14/2021		

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	06/14/2021				

## Labeler - Shimmy Products LLC (117838009)

Revised: 6/2021 Shimmy Products LLC