## SANIZEN- alcohol gel Village Pharma, 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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#### Sanizen Gel

#### **DRUG FACTS:**

#### **Active Ingredients:**

Ethyl Alcohol v/v 75%

Antiseptic

#### Uses:

Hand sanitizer to help reduce bacteria on the skin.

### **Warnings:**

For external use only.

Flammable. Keep away from fire or flame.

## When using this product,

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

### Stop use and ask a doctor if

irritation or rash appears and lasts.

## Keep out of reach of children.

Children must be supervised in the use of this product. If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions:**

Put enough product in your palm to cover hands and rub hands together briskly until dry.

#### Other Information:

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces.

#### **Inactive Ingredients:**

Water (Aqua), Aloe Barbadensis Leaf Extract, Vitamin E, Carbomer, Triethanolamine, DMDM Hydantoin, Glycerol

#### **Package Labeling:**



DRUG FACTS: Active Ingredients: Antiseptic Ethyl Alcohol v/v Uses: Hand sanitizer to help reduce bacteria on the skin. Warnings: For external use only. Flammable. Keep away from fire or flame. When using this product, do not use in or near the eyes. In case of contact, rinse eyes Stop use and ask a doctor if irritation or rash appears and lasts. Keep out of reach of children. Children must be supervised in the use of this product. If swallowed, get medical help or contact a Poison Control Center right away. Directions:
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Manufactured for: Village Pharma, LLC Agoura Hills, CA 91301 Questions or Comments? Please email: info@villagepharma.com Made in China



## **SANIZEN**

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71574-711
Route of Administration	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)		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
.ALPHATOCOPHEROL (UNII: H4N855PNZ1)		
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)		
TROLAMINE (UNII: 9O3K93S3TK)		
DMDM HYDANTO IN (UNII: BYR0546 TOW)		
GLYCERIN (UNII: PDC6A3C0OX)		

l	Packaging				
l	#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
l	1	NDC:71574-711-10	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/13/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/13/2020	

# Labeler - Village Pharma, LLC (080749749)

Revised: 4/2020 Village Pharma, LLC