

## **HAND SANITIZER- ethyl alcohol gel**

### **Medustry 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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#### **Active Ingredient**

Ethyl Alcohol 75%

#### **Purpose**

Antiseptic

#### **Uses**

For hand washing to decrease bacteria on the skin.

Recommended for repeated use.

#### **Warnings**

Flammable. Keep away from fire or flame.

For external use only.

Avoid contact with broken skin.

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. If swallowed, get medical help.

#### **Directions**

Not recommended for infants.

Wet hands thoroughly with product and allow to dry without wiping.

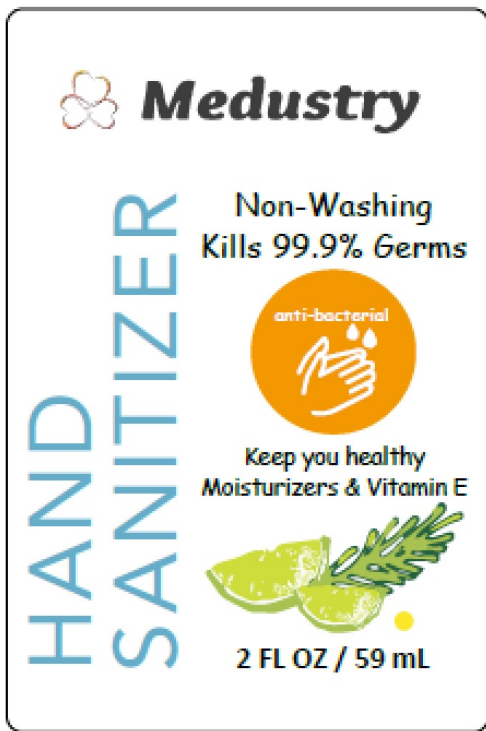
Children under 6 years of age should be supervised when using this product.

#### **Other Information**

Do not store above 110°F. May discolor some fabrics.

#### **Inactive Ingredients**

Water, Aloe Barbadensis Leaf Juice, Maltodextrin, Glycerin, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate crosspolymer, Triethanoamine, Fragrance, Tocopheryl Acetate.



## HAND SANITIZER

ethyl alcohol gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74418-202
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
MALTO DEXTRIN (UNII: 7CVR7L4A2D)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
TROLAMINE (UNII: 9O3K93S3TK)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74418-202-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2020	
<b>Marketing Information</b>				
<b>Marketing Category</b>		<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph not final		part333E	04/24/2020	

**Labeler** - Meduistry LLC (117306610)

Revised: 4/2020

Meduistry LLC