

**HAND SANITIZER- alcohol gel**  
**In Designs Global 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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**Hand Sanitizer**

***Drug Facts***

***Active ingredient***

Alcohol 62%

***Purpose***

Antiseptic

***Use***

- For handwashing to decrease bacteria on the skin

***Warnings***

**For external use only**

- **Flammable**, keep away from fire and flame
- Does not contain grain alcohol; do not drink. If taken internally serious gastric disturbances will result.

**When using this product**

- Avoid the eyes and mucous membranes
- In the case of eyes or mucous membranes contact, rinse area thoroughly with water
- In case of deep or puncture wounds or serious burns, consult physician.
- If redness, irritation, swelling, or pain persists or increases or if infection occurs discontinue use and physician.

**Keep out of reach of children**

- If swallowed, contact a doctor or Poison Control Center immediately.

***Directions***

- Rub dime sized amount between hands until dry.
- Supervise children in the use of this product.
- In the case of eye contact, rinse eyes thoroughly with water.

***Other information***

- Store below 105 °F
- May discolor some fabrics.

***Inactive ingredients***

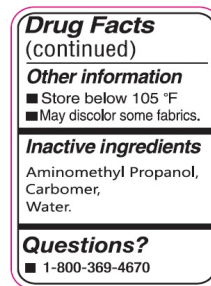
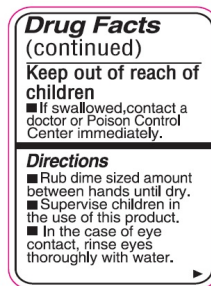
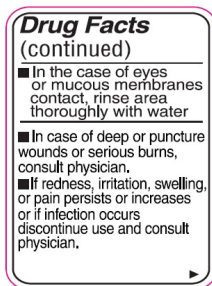
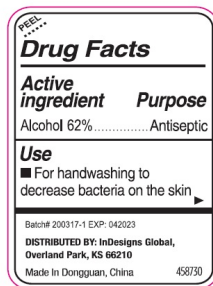
Aminomethyl Propanol, Carbomer, Water.

## Questions?

- 1-800-369-4670

## Package Labeling:

35x47mm



## HAND SANITIZER

alcohol gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74712-000
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
WATER (UNII: 059QF0K00R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74712-000-01	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/10/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/10/2020	

