# HAND SANITIZER PREMIUM FORMULA- hand sanitizer gel LTP Global

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(s)**

Alcohol 70% v/v. Purpose: Antiseptic

### **Purpose**

Antiseptic

#### Use

- to decrease bacteria that potentially can cause disease
- recommended for repeated use

# Warnings

For external use only: hands

Flammable. Keep away from heat or flame.

- When using this product keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommend for infacts

#### Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)
- Do not use on children under two months of age.

#### **Inactive ingredients**

Water, Glycerin, Carbomer, Trolamine, L-Menthol, Green Tea Leaf, Aloe Vera Leaf

#### Package Label - Principal Display Panel

# the simple market

NDC 73812-002-30

# HAND SANITIZER PREMIUM FORMULA

Ethanol 70 %

Leaves hands soft with aloe extract

Easily sterilize harmful bacteria without water or soap

Quick drying, refreshing, leaving non-sticky hands









### HAND SANITIZER PREMIUM FORMULA

hand sanitizer gel

# **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73812-002

Route of Administration TOPICAL

### Active Ingredient/Active Moiety

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Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL		

Strength

Packaging					
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:73812-002-	300 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/20/2020		

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	05/20/2020				

# Labeler - LTP Global (694215916)

Establishment				
Name	Address	ID/FEI	Business Operations	
ICOMAX Co., Ltd.		694834031	manufacture(73812-002)	

Revised: 6/2020 LTP Global