# GERM FREE DISINFECTANT GEL- hand sanitizer gel Guangzhou Mengfeishi Pharmaceutical Technology Co., Ltd.

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**

Ethanol 70% (v/v)

#### **Purpose**

Antibacterial sterilization

#### Use

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

# Warnings

For external use only, Flammable. Keep away from heat of flame

#### Do not use

In 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 occurs. These may be signs of asrtious condition.

#### Keep out of reach of children.

If swallowed, get medical help or contact a poison Control Center right away.

#### **Directions**

Take appropriate amount of this product, rub with both hands until dry, no need to wash by hand.

#### Other information

Store between 15-30C (59-86F)

Avoid freexing and excessive heat above 40C(104F)

#### **Inactive ingredients**

water, propanediol, carbomer, triethanolamine, chlorphenesin, aloe barbadensis leaf extract.

### Questions?

+83 20 36632232

# Package Label - Principal Display Panel



# GERM FREE DISINFECTANT GEL

hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77311-016
Route of Administration	TOPICAL		

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

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
PROPANEDIOL (UNII: 5965N8W85T)			
CARBO MER HO MO PO LYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)			
TROLAMINE (UNII: 9O3K93S3TK)			
CHLO RPHENESIN (UNII: 1670 DAL4SZ)			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)			

Packaging					
l	#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
l	1	NDC:77311-016-02	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/10/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	06/10/2020		

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Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Guangzhou Mengfeishi Pharmaceutical Technology Co., Ltd.		402685342	manufacture(77311-016)	

Revised: 5/2020

Guangzhou Mengfeishi Pharmaceutical Technology Co., Ltd.