

GERM FREE INSTANT HAND SANITIZER- 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.

Main active ingredients and content

This product contains 75% ± 5% ethanol

Product description

Clean and bacteriostatic, inhibit staphylococcus aureas, escherichia coli, candida albicans. Prevent bacterial growth

Inhibition of microorganisms

This product has a strong and effective inhibitory effect on staphylococcus aureus, escherichia coli and candidaalbicans

Dosage form

Gel

Usage

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

Note

This product is an external product and should not be taken orally. In case of accidental contact with eyes, please rinse immediately with plenty of water or seek medical advice. Avoid open flames.

Keep out of reach of children

Keep out of reach of children.

Main ingredients

Water, ethanol, propanediol, carbomer, triethanolamine, chlorphenylglycerin, ALOE BARBADENSIS leaf extract

Storage

Seal and store in a cool place

Package Label - Principal Display Panel

NDC 77311-500-02

Product description: Clean and bacteriostatic, inhibit staphylococcus aureus, escherichia coli, candida albicans. Prevent bacterial growth

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Made in China



75%Alcohol

INSTANT HAND SANITIZER
Kills up to 99.99% of Germs

Hygiene license of production enterprise: ywxxx (20171-01--no 8003
Executive standard: GB/T 26373-2010 qualified
Manufacturer: Guangzhou mengleishi Pharmaceutical Technology Co., Ltd
Production address: No.1, community 14, Luogang, JUNHE street, Baiyun District, Guangzhou



MFS:17/04/2020
EXP:16/04/2022



Blue Mineral Corporation
bluemineral.com

16.9 FL. OZ. - 500mL

77311-500-03



75%
Alcohol

INSTANT HAND SANITIZER

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GERM FREE INSTANT HAND SANITIZER

hand sanitizer gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77311-500
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
WATER (UNII: 059QF0K00R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPANEDIOL (UNII: 5965N8W85T)	
TROLAMINE (UNII: 9O3K93S3TK)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77311-500-02	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/20/2020	
2	NDC:77311-500-03	500 mL in 1 BOTTLE; 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 - Guangzhou Mengfeishi Pharmaceutical Technology Co., Ltd. (402685342)**Establishment**

Name	Address	ID/FEI	Business Operations
Guangzhou Mengfeishi Pharmaceutical Technology Co., Ltd.		402685342	manufacture(77311-500)

Revised: 5/2020

Guangzhou Mengfeishi Pharmaceutical Technology Co., Ltd.