INSTANT HAND SANITIZER- instant hand sanitizer gel FOSHAN KANGKANG BIOTECHNOLOGY?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.

INSTANT HAND SANITIZER

Active Ingredient(s)

Alcohol 71% v/v

Purpose

Antimicrobial

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease

Warnings

For external use only; Do not eat ;Flammable,keep away from fire

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 a serious condition.

keep out of reach of children

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

Directions

Place enough product on hands to cover all surfaces. Rub hands together until dry.

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

Other information

Inactive Ingredients

Glycerin







Active Ingredient(s): Alcohol71% v/v

Purpose:Antimicrobial

Use:Hand Sanitizer to help reduce bacteria that potentially can cause disease

Warnings:For external use only; Do not eat; Flammable, keep away from fire;

Do not use:In children less than 2 months of age; On open skin wounds

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MADE IN CHINA

Otc-stop use section:Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition

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INSTANT HAND SANITIZER

instant hand sanitizer gel

Product Information

HUMAN OTC DRUG NDC:78413-001 Product Type Item Code (Source)

TOPICAL **Route of Administration**

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ALCOHOL ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) $71\,mL\,$ in $100\,mL\,$

Inactive Ingredients

Ingredient Name Strength

GLYCERIN (UNII: PDC6A3C0OX)

Packaging

1	# Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78413-001-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/09/2020	
2	NDC:78413-001-02	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/09/2020	
3	NDC:78413-001-03	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/09/2020	

Marketing Information

Marketing Category	Marketing End Date		
OTC monograph not final	part333A	06/09/2020	

Labeler - FOSHAN KANGKANG BIOTECHNOLOGY?CO.,LTD (554534671)

Registrant - FOSHAN KANGKANG BIOTECHNOLOGY?CO.,LTD (554534671)

Establishment							
Name	Address	ID/FEI	Business Operations				
FOSHAN KANGKANG BIOTECHNOLOGY?CO.,LTD		554534671	manufacture(78413-001)				

Revised: 6/2020 FOSHAN KANGKANG BIOTECHNOLOGY?CO.,LTD