INSTANT HAND SANITIZER- alcohol gel
Zhejiang Bayside Biotech 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.

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antimicrobial

Purpose

Antimicrobial, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria on the skin.

Warnings

- For external use only. Keep out of reach of children.
- Flammable. Keep away from source of heat or fire.

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product avoid contact with eyes. If contact occurs in case, rinse with water.

Stop using this product and ask doctor if irritation or redness develops and lasts.

Keep out of reach of children.

Directions

- Wet hands thoroughly and rub together until dry.
- Children under 6 years of age should be supervised when using instant hand sanitizer.

Other information

- Seal and store at a temperature below 40C (104F)

Inactive ingredients

glycerin, water, Isopropyl Myristate, Carbomer, Aloe Leaf juice, Tocopheryl Acetate, Aminomethyl propanol, Rose oil.

Package Label - Principal Display Panel

266 mL NDC: 78766-001-01

Kills more than 99.99% of most common germs that may make you sick

Drug facts

Active ingredient **Purpose**
Ethyl alcohol 70% v/v.....Antimicrobial

Inactive ingredients
Water(Aqua), Glycerin, Isopropyl Myristate, Carbomer, Purified Inner Leaf Fillet Juice, Tocopheryl Acetate, Aminomethyl Propanol, Rose essential oil .

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3.When using this product avoid contact with eyes. If contact occurs in case , rinse with water.
4.Stop using this product and ask doctor if irritation or redness develops and lasts.

Storage
Seal and Store at a temperature below 40℃.

FDA Registration No.:3016735726
Question or comments? 1 877 865 6938
Distributed by:Zee Scientific LLC
One International BLVD.Mahwah,Suite 400, NJ 0743
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INSTANT HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78766-001
Route of Administration	EXTRACORPOREAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	3 in 100 mL
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	0.5 in 100 mL
WATER (UNII: 059QF0KO0R)	25.9 in 100 mL
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	0.04 in 100 mL
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	0.05 in 100 mL
ALOE FEROX LEAF (UNII: 0D145J8EME)	0.1 in 100 mL
CARBOMER 940 (UNII: 4Q93RCW27E)	0.4 in 100 mL
ROSE OIL (UNII: WUB68Y35M7)	0.01 in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78766-001-01	266 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
2	NDC:78766-001-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/19/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/13/2020	

Labeler - Zhejiang Bayside Biotech Co., Ltd. (554477072)

Registrant - Zhejiang Bayside Biotech Co., Ltd. (554477072)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Bayside Biotech Co., Ltd.		554477072	manufacture(78766-001) , label(78766-001)

Revised: 6/2020 Zhejiang Bayside Biotech Co., Ltd.