EGOLAN INSTANT HAND SANITIZER GEL- ethyl alcohol gel Sanrace 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.

EGOLAN[®] Instant Hand Sanitizer gel

Drug Facts

Active Ingredient Ethyl Alcohol 75% v/v

Purpose

Antiseptic

Use[s]

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 or 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 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.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-83F)
- Avoid freezing and excessive heat above 40C (104F)

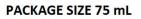
Inactive Ingredients:

Water, Glycerin, Propylene glycol, Carbomer, Aloe yohjyu matsu ekisu, Triethanolamine, Disodium Edta

*99.99% of most common germs.

Manufacturer: Sanrace Biotechnology Co.,Ltd Manufacturer Address:268 Yanzhou Road, Lanxi Development Zone,Zhejiang Province. Made In China

Packaging





PACKAGE SIZE 3785 mL





1 Gal/3785ml

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etl	ıyl alcohol gel									
P	roduct Informa	tion								
P	roduct T ype		HUMAN OTC DRUG	Ite m Co	m Code (Source)		NDC:75448-031			
R	oute of Administra	ation	TOPICAL							
Active Ingredient/Active Moiety										
Ingredient Name					Basis of Strength		Strength			
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)					ALCOHOL	nL in 100 mL				
Inactive Ingredients										
			Ingredient Name				Strength			
w	ATER (UNII: 059QF	0 KO 0 R)								
GLYCERIN (UNII: PDC6A3C0OX)										
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)										
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)										
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)										
TROLAMINE (UNII: 903K93S3TK)										
EDETATE DISODIUM ANHYDROUS (UNII: 8 NLQ36 F6 MM)										
P	ackaging									
#	Item Code		Package Description		Marketing Start Date	Mark	eting End Date			
1	NDC:75448-031-02		LE; Type 0: Not a Combination 1	Product	01/05/2021		0			
2	NDC:75448-031-01	3785 mL in 1 BO	TTLE; Type 0: Not a Combinatio	n Product	0 1/0 5/20 21					
Marketing Information										
U			on Number or Monograph C	Citation	Marketing Start Date	Marketing End Date				
			5.		0 1/0 5/20 21		-			
	-									

Labeler - Sanrace Biotechnology Co., Ltd. (543000938)

Establishment									
Name	Address	ID/FEI	Business Operations						
Sanrace Biotechnology Co., Ltd.		543000938	manufacture(75448-031)						

Revised: 1/2021

Sanrace Biotechnology Co., Ltd.