EGOLAN- 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® HAND SANITIZER GEL

DRUG FACTS

Active ingredient

Ethyl Alcohol 75%

Purpose

Antiseptic

Use: Hand sanitizer to help reduce bacteria on the skin.

Warnings

Flammable. Keep away from fire or flame.

For external use only. do not inhale or ingest.

Do not use in, or near the eyes. In case of contact with eyes, rinse thoroughly with water.

Avoid contact with broken skin

Stop use and ask a doctor if irritation or rash develops and persists for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- Wet hands thoroughly with product and allow to dry without wiping.
- For children under 6, use only under adult supervision
- Not recommended for infants

Other Information

- Store below 110°F (43°C)
- May discolor certain fabrics or surfaces

Inactive ingredients

Water,Glycerin,Propylene Glycol,Triethanolamine,Carbomer,Aloe Vera Gel

ANTIBACTERIAL

KILLS 99.9% GERMS

Manufacturer:

Sanrace Biotechnology Co., Ltd

Manufacturer Address:268 Yanzhou Road,

Lanxi Development Zone,Zhejiang Province.

Made In China.

Packaging

EGOLAN

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75448-023
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: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:75448-023-03	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/2020	

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

Establishment

Name	Address	ID/FEI	Business Operations
Sanrace Biotechnology Co., Ltd.		543000938	manufacture(75448-023)

Revised: 6/2020 Sanrace Biotechnology Co., Ltd.