EUNOS- alcohol gel
YIRONG TRADING (NANJING) 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

Ethyl Alcohol 75% v/v.

Purpose

Antimicrobial

Uses

Hand Sanitizer to help reduce bacteria on the skin that potentially can cause disease.

Warnings

Flammable. Keep away from fire or flame.

For external use only.

When using this product

- do not use in or near the eyes. In case of contact with eyes, rinse eyes thoroughly with water.
- avoid using on broken skin.

Stop use and ask a doctor if irritation or rash appears and lasts.

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

Directions

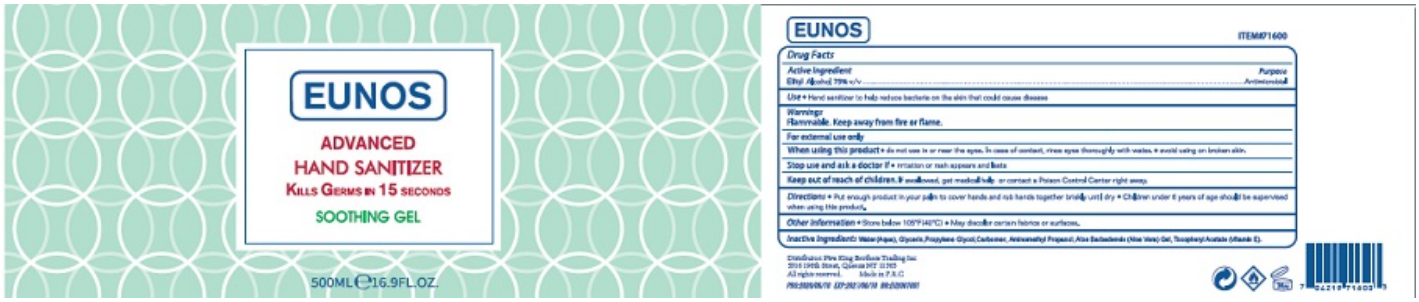
- Put enough product in your palm to cover hands and rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using this product.

Inactive ingredients

Aqua, Glycerin, Propylene Glycol, Carbomer, Aminomethyl Propanol, Aloe Barbadensis (Aloe Vera) Gel, Tocopheryl Acetate (Vitamin E)

Package Label - Principal Display Panel

500 mL Label



1000 mL Label

EUNOS

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75007-002
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)	
CARBOMER HOMOPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: F68VH75CJC)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75007-002-01	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2020	
2	NDC:75007-002-02	1000 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 - YIRONG TRADING (NANJING) CO., LTD (554529568)

Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Meizhiyuan Cosmetics Co., Ltd		543036597	manufacture(75007-002)

Revised: 6/2020 YIRONG TRADING (NANJING) CO., LTD