HAND SANITIZER- ethyl alcohol gel
Yuyao Jessie Commodity 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.

51414-912 HAND SANITIZER Ethyl Alcohol 62%

Active Ingredient(s)

Ethyl Alcohol 62%

Purpose

Antiseptic

Use

for hand sanitizing

Warnings

For external use only.
Flammable, Keep away from heat or flame,
When using this product keep out of eyes, flush thoroughly with water. Avoid contact with broken skin. Do not inhale or ingest.

Stop use and ask a doctor if irritation develops.

Keep out of reach of children. except under adult supervision.

If swallowed, get medical help or call a poison control center right away.

Directions

Wet hands thoroughly with product and allow to dry without wiping.

For children under 6 age use only under adult supervision.

Not recommended for infants.

Inactive ingredients

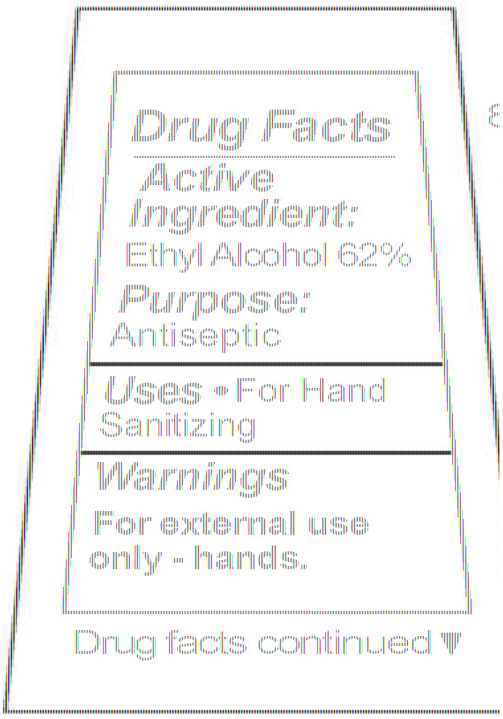
Carborner,Fragrance,Glycerin,Propylene Glycol,TocopherylAcetate,Triethanolamine,Water.

Package Label - Principal Display Panel

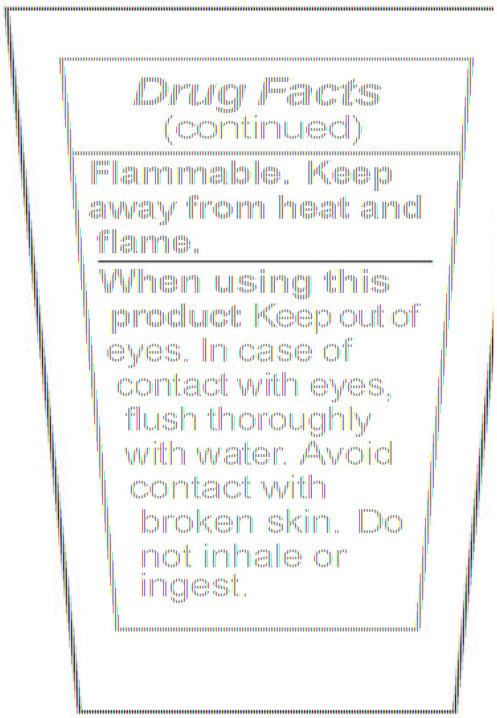
Item:9054

L*H=25*45MM

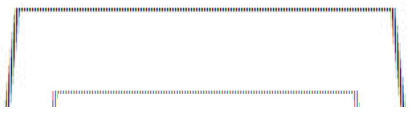
first layer-front side
第一层正面



first layer-back side
第一层背面



second layer-front side
第二层正面



第二层正面

Stop use and ask a doctor if skin irritation develops.

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

Directions

- Wet hands thoroughly with product and allow to dry without wiping.

Drug facts continued ▼

second layer-back side
第二层背面

Drug Facts
(continued)

- For children under 6 age use only under adult supervision.
- Not recommended for infants.

Other Information

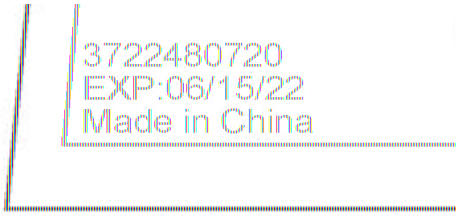
- Do not store above 105°F.
- May discolor some fabrics.

third layer-front side
第三层正面

Inactive Ingredients

Carbomer, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate, Triethanolamine, Water.

1oz 30ML



HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51414-912
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51414-912-01	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
2	NDC:51414-912-02	10 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
3	NDC:51414-912-03	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
4	NDC:51414-912-04	20 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
5	NDC:51414-912-05	480 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
6	NDC:51414-912-06	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
7	NDC:51414-912-07	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
8	NDC:51414-912-08	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	

9	NDC:51414-912-09	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
10	NDC:51414-912-10	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
11	NDC:51414-912-11	90 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
12	NDC:51414-912-12	8 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
13	NDC:51414-912-13	0.9 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
14	NDC:51414-912-14	5 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
15	NDC:51414-912-15	25 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
16	NDC:51414-912-16	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
17	NDC:51414-912-17	35 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
18	NDC:51414-912-18	75 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	
19	NDC:51414-912-19	18 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/10/2020	

Labeler - Yuyao Jessie Commodity Co., Ltd. (529892305)

Registrant - Yuyao Jessie Commodity Co., Ltd. (529892305)

Establishment

Name	Address	ID/FEI	Business Operations
Yuyao Jessie Commodity Co., Ltd.		529892305	manufacture(51414-912)

Revised: 3/2022 Yuyao Jessie Commodity Co., Ltd.