

KANGKANG ANTISEPTIC HAND SANITIZER ETHYL ALCOHOL 75- alcohol liquid
Jiangsu Unicorn Electronic Technology 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.

KANGKANG Antiseptic Hand Sanitizer Spray Ethyl Alcohol 75%

Drug Facts

Active ingredient

Ethyl alcohol 75%

Purpose

Antiseptic

Use

For hand washing to decrease bacteria on the skin

Warnings

For external use only.

Flammable, keep away from fire or flame

Do not use

in the eyes

Stop use and ask a doctor if:

- irritation and redness develop
- condition persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

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

Inactive ingredients

Water.

Package Labeling:



Drug Facts	
Active ingredient Ethyl alcohol 75%	Purpose Antiseptic
Use For hand washing to decrease bacteria on the skin	
Warnings For external use only. Flammable, keep away from fire or flame Do not use in the eyes Stop use and ask a doctor if: ■ irritation and redness develop ■ condition persists for more than 72 hours	
DISTRIBUTED BY JIANGSU UNICORN ELECTRONIC TECHNOLOGY CO., LTD. F2, NO.9 SHIDAI AVENUE, QUANSHAN DISTRICT, XUZHOU, JIANGSU, CHINA MADE IN CHINA	
TO REPORT A SERIOUS ADVERSE EVENT, CONTACT: 909-323-9867	
 6 973 146 070 029	

Drug Facts (continued)
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
Directions Wet hands thoroughly with product and allow to dry without wiping.
Inactive ingredients Water.

KANGKANG ANTISEPTIC HAND SANITIZER ETHYL ALCOHOL 75

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:90074-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.75 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:90074-000-01	13 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	part333E	10/10/2020	

