

HAND SANITIZER 1.8 OZ ANTISEPTIC- ethyl alcohol liquid
China Ningbo Shangge 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.

Hand Sanitizer Spray 1.8oz Antiseptic

Ethyl Alcohol 70.0% v/v

Antiseptic

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

For external use only: hands

Flammable, keep away from fire or flame

- keep out eyes. In case of contact with eyes, flush thoroughly with water
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.

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

- spray hand thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

- store at room temperature
- may discolor certain fabrics
- harmful to wood finished and plastics

water, glycerin, tocopheryl acetate, aloe barbadensis leaf juice, fragrance.

XtraCare®

Kills 99.9%
of Germs*

Hand Sanitizer

With
moisturizers
& vitamin E

On-The-Go
Purse or
Pocket Size

CAUTION:
EYE IRRITANT
Read back label before use.

2 FL OZ (59 mL)

XtraCare®

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Hand Sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 70.0% v/v.....

Purpose

Antiseptic

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Other information

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Questions/Comments? 1-855-345-5575

Inactive ingredients water, glycerin, tocopheryl acetate, aloe barbadensis leaf juice, fragrance.

*Effective at eliminating 99.9% of many common harmful germs and bacteria.

DISTRIBUTED BY: REJOICE INTERNATIONAL CORP.
NORTHVILLE, MI 48167, USA
MADE IN CHINA



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With moisturizers & vitamin E



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MADE IN CHINA

HAND SANITIZER 1.8 OZ ANTISEPTIC

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58503-123
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:58503-123-02	1 in 1 BLISTER PACK	03/05/2020	
1	NDC:58503-123-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/05/2020	

Labeler - China Ningbo Shangge Technology Co., Ltd. (529287434)

Establishment

Name	Address	ID/FEI	Business Operations
China Ningbo Shangge Technology Co., Ltd.		529287434	manufacture(58503-123)

Revised: 3/2020

China Ningbo Shangge Technology Co., Ltd.