

HAND SANITIZER 2PK ALOE 2OZ- 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.

Active Ingredient(s)

Ethyl Alcohol 70% v/v.

Purpose

Antimicrobial

Use

- for handwashing to decrease bacteria on the skin
- recommended for repeated use

Warnings

For external use only. Flammable, keep away from heat and flame

Do not use

- in the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or 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.

Directions

- Wet hands thoroughly with product
- briskly rub hands together until dry.
- Supervise children under 6 years of age when using this product.

Other information

- Store at room temperature
- may discolor certain fabrics

Inactive ingredients

water, carbomer, triethanolamine, glycerin, propylene glycol, fragrance, aloe barbadensis leaf juice, tocopheryl acetate, fd&c blue no. 1, fd&c yellow no.5.

1-855-345-5575

Package Label - Principal Display Panel

XtraCare®

70% ALCOHOL

**HAND
Sanitizer**
ANTIBACTERIAL

Moisturizes with
Vitamin E & Aloe



2 FL OZ (59 mL)

Drug Facts

Active ingredient

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Directions • wet hands thoroughly with product • briskly rub hands together until dry • supervise children under 6 years in the use of this product

Other information

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Inactive ingredients water, carbomer, triethanolamine, glycerin, propylene glycol, fragrance, aloe barbadensis leaf juice, tocopheryl acetate, fd&c blue no.1, fd&c yellow no.5.

Questions/comments? 1-855-345-5575

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



HAND SANITIZER 2PK ALOE 2OZ

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58 503-152
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
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0K00R)	
TROLAMINE (UNII: 9O3K93S3TK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	

ALOE (UNII: V5VD430YW9)

CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)

ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

FD&C YELLOW NO. 5 (UNII: I753WB2F1M)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58503-152-01	1 in 1 CONTAINER	09/28/2020	
1		118 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	part333A	09/28/2020	

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

Establishment

Name	Address	ID/FEI	Business Operations
China Ningbo Shangge Technology Co., Ltd.		529287434	manufacture(58503-152)

Revised: 9/2020

China Ningbo Shangge Technology Co., Ltd.