

**HAND SANITIZER- hand sanitizer liquid**  
**Fortress Expert 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.*

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**73549-026**  
**HAND SANITIZER**

***Active ingredient***

Ethyl Alcohol 70%

**Purpose**

Antiseptic

***Use***

■ For handwashing to decrease bacteria on the skin.

***Warnings***

For external use only: hands  
Flammable, keep away from fire or flame

**When using this product**

- keep out of 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
- condition persists for more than 72 hours

**Keep out of reach of children**

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

- for children under 6, use only under adult supervision
- not recommended for infants

### Other information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

### Inactive ingredients

Water(Aqua), PEG-40 Hydrogenated Castor Oil, Fragrance(Parfum) , Acrylates/ C10-30Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Aloe Barbadensis Extract, Butylene Glycol, 1,2-Hexanediol, Caprylhydroxamic Acid, Citric Acid, Disodium EDTA, Tocopheryl Acetate, Red 33, Yellow 5

### Package Label

73549-026-01 30ml



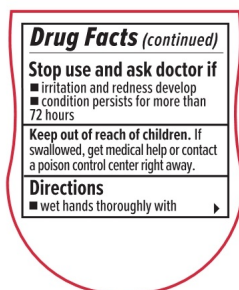
30x36mm



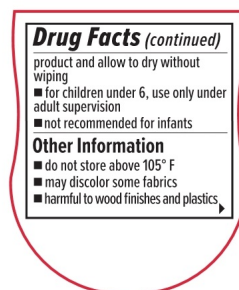
outside front panel  
**1**



inside left flap  
**2**



inside center panel  
**3**



inside right flap  
**4**



outside right panel  
**5**

## HAND SANITIZER

hand sanitizer liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:73549-026
<b>Route of Administration</b>	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 YELLOW NO. 5 (UNII: I753WB2F1M)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ALOE VERA WHOLE (UNII: KIZ4X2EHYX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FRAGRANCE FLORAL ORC0902236 (UNII: R66Z4YW3X0)	
WATER (UNII: 059QF0KO0R)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73549-026-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/30/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/30/2020	

**Labeler** - Fortress Expert Co., Ltd (543358697)**Establishment**

Name	Address	ID/FEI	Business Operations
Fortress Expert Co., Ltd		543358697	manufacture(73549-026)

Revised: 6/2021

Fortress Expert Co., Ltd