

SMART CARE HAND SANITIZER 2 OZ- ethyl alcohol gel
Guangzhou Xujohn Bio-Technique 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.

Smart Care® Hand SANITIZER

Drug Facts

Active Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Use

To help reduce bacteria and germs on the skin.

WARNING

Flammable. Keep away from fire or flame. For external use only

- Stop use and ask a doctor if irritation or redness develops and persists.
- **Keep out of reach of children.** • In case of accidental digestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

- Place enough product in palm to cover hands and rub hands together briskly until dry.
- Children under 6, use only under adult supervision.
- Not recommended for infants.

Other Information

- Do not store above 100° F (38°C).
- May discolor some fabrics.
- Harmful to wood finishes and plastics.

Inactive Ingredients

WATER(AQUA), GLYCERIN, ACRYLATES/C10-30 ALKYL ACRYLATE
CROSSPOLYMER, AMINOMETHYLPROPANOL, TOCOPHERYL ACETATE, ALOE
BARBADENSIS LEAF JUICE, MALTODEXTRIN, POTASSIUM SORBATE, SODIUM
BENZOATE, CI 42090, CI 19140

QUESTIONS OR COMMENTS?

1-877-274-8358 Toll Free in USA

1-909-434-0911 International

KILLS UP TO 99.9% OF MOST COMMON GERMS

Aloe Vera & Vitamin E

FAST & EFFECTIVE

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DESIGNED IN U.S.A. • MADE IN CHINA

Packaging



SMART CARE HAND SANITIZER 2 OZ

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75351-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75351-001-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/06/2020	

Labeler - Guangzhou Xujohn Bio-Technique Co., Ltd. (403308789)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Xujohn Bio-Technique Co., Ltd.		403308789	manufacture(75351-001)

Revised: 4/2020

Guangzhou Xujohn Bio-Technique Co., Ltd.