

SMART CARE HAND SANITIZER 2 OZ- ethyl alcohol gel
Shenzhen Lantern Science 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), ISOPROPYL ALCOHOL, GLYCERIN, CARBOMER, AMINOMETHYL PROPANOL, PARFUM, PROPYLENE GLYCOL, ISOPROPYL MYRISTATE, ALOE BARBADENSIS LEAF JUICE, TOCOPHERYL ACETATE, FD&C YELLOW NO. 5 (TARTRAZINE), FD&C BLUE NO. 1

KILLS UP TO 99.99% OF MOST COMMON GERMS

Aloe Vera & Vitamin E

FAST & EFFECTIVE

DESIGNED IN U.S.A. • MADE IN CHINA

QUESTIONS OR COMMENTS?

1-877-274-8358 Toll Free in USA. • 1-909-434-0911 International

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DISTRIBUTED BY ASHTEL STUDIOS INC. ONTARIO, CALIFORNIA 91761

Packaging



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QUESTIONS OR COMMENTS? ITEM #: 92312-24
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SMART CARE HAND SANITIZER 2 OZ
ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-192	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL	
Inactive Ingredients				
	Ingredient Name		Strength	
	WATER (UNII: 059QF0KO0R)			
	ISOPROPYL ALCOHOL (UNII: ND2M416302)			
	GLYCERIN (UNII: PDC6A3C0OX)			
	CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
	AMINOMETHYLPROPANOL (UNII: LU49E6626Q)			
	PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
	ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)			
	ALOE VERA LEAF (UNII: ZY81Z83H0X)			
	.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)			
	FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
	FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-192-01	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 - Shenzhen Lantern Science Co., Ltd. (421222423)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co., Ltd.		421222423	manufacture(54860-192)

Revised: 4/2020

Shenzhen Lantern Science Co., Ltd.