

SMART CARE HAND SANITIZER 8 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 Ingredients

Ethyl Alcohol 62%

Purposes

Antiseptic

Uses

For external use only: hands

Directions:

Put a thumbnail size amount in your palm and rub your hands together briskly until dry. Children under 6 years of age should be supervised when using SMART CARE® HAND SANITIZER. 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

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.

Keep out of reach of children

If swallowed get medical help or contact a Poison Control Center right away.

KILLS UP TO 99.99% OF MOST COMMON GERMS

Aloe Vera & Vitamin E

FAST & EFFECTIVE

SMARTCAREUS.COM

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

DESIGNED IN U.S.A.

MADE IN CHINA

Packaging



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 50016 92311 0 8	
ITEM #: 92311-12	

SMART CARE HAND SANITIZER 8 OZ

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-190
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 ACETATE (UNII: 9E8X80D2L0)	
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:54860-190-01	236.6 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(54860-190)

