SMART CARE LUXURY HAND SANITIZER 1.35OZ - GOLDEN GROVE- alcohol spray
Shenzhen Lantern Scicence Co.,Ltd.

Smart Care Luxury Hand Sanitizer Spray 1.35oz - Golden Grove

Drug Facts

Active Ingredient

Active ingredient Purpose

Ethyl Alcohol 70% v/v Antiseptic

Uses

Decreases bacieria on skin

Warning

For external use only

Flammable, keep away from fire or flame.

When using this product

Do not get into eyes. lf contact occurs, rinse thoroughly with water.

Do not use

lf you are allergic to any of the ingredients

Stop use and ask a doctor

lf irritation or rash develops and continues for more than72 hours.

keep out of reach of children

lf swallowed, get medical help or contact a Poison Control Center immediately.

Directions

Adults and children 6 years and older
Spray onto hands and rub together unil dry. Recommended for repeat use.
Children under 6 years of age

should be supervised when using this product.

Inactive ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Gel, Aminomethyl Propanol,Buiue 1, Denatonium Benzoate, Fragrance, Glycerin, Peg-40 Hydrogenated Castor Oil, Red 4, Tocopheryl Acetate, Water, Yellow 5.

other Information

Storebetween15-30C(59-86F)
Avoid freezing and excessive heat above 40C (104F)

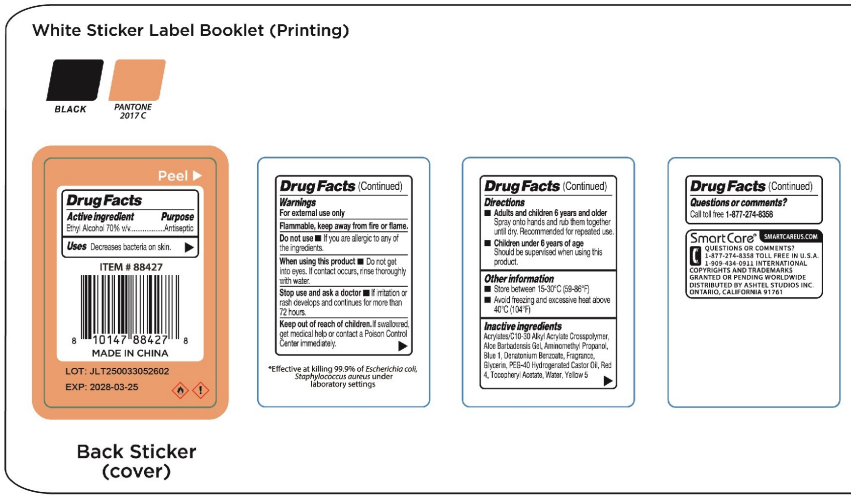
Spray 3-4 times each time.
Used in the skin area of the hands.

packing

Packaging

Front View Back View

Product dimensions: (W)58mm x (H)104mm x (D)17mm



Back Sticker (cover)

SMART CARE LUXURY HAND SANITIZER 1.35OZ - GOLDEN GROVE

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54860-463
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)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-463-01	40 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/09/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	04/09/2025	

Labeler - Shenzhen Lantern Scicence Co.,Ltd. (421222423)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-463)

Revised: 4/2025 Shenzhen Lantern Scicence Co.,Ltd.