

SMART CARE SWEET PEA HAND SANITIZER SWEET PEA 01- alcohol spray
Shenzhen Lantern Science Co.,Ltd.

88398 Smart Care Sweet Pea Hand Sanitizer Spray sweet pea

Drug Facts

Active Ingredient

Active ingredient Purpose

Ethyl Alcohol 70% v/v Antiseptic

Uses

Decreases bacteria on skin

Warning

For external use only

Do not use

If you are allergic to any of the ingredients

When using this product

Do not get into eyes. If contact occurs, rinse thoroughly with water. Do not use in or near eyes. In case of eye contact, flush eyes thoroughly with water. Discontinue if skin becomes irritated and ask for a doctor.

Stop use and ask a doctor

If irritation or rash develops and continues for more than 72 hours.

Keep out of reach of children

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

Inactive ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Gel, Aminomethyl Propanol, Blue 1, Denatonium Benzoate, Fragrance, Glycerin, PEG-40 Hydrogenated Castor Oil, Red 4, Red 33, Tocopheryl Acetate, Water

Directions

Adults and children 6 years and older

Spray onto hands and rub together until dry. Recommended for repeat use.

Children under 6 years of age

should be supervised when using this product.

other Information

Store between 15-30C (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



SMART CARE SWEET PEA HAND SANITIZER SWEET PEA 01
alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-450	
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	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)				
FD&C RED NO. 4 (UNII: X3W0AM1JLX)				
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)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-450-01	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/16/2025	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003		03/16/2025	

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

Establishment			
Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-450)