

**SMART CARE HAND SANITIZER WITH SILICONE CASE 3PK- alcohol spray**  
**Shenzhen Lantern Science Co.,Ltd.**

-----  
**Smart Care Hand Sanitizer with Silicone Case 3pk**

**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.

**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.

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Gel, Aminomethyl Propanol, Denatonium Benzoate, Fragrance, Glycerin, Peg-40 Hydrogenated Castor Oil, Red 4, Red 33, Tocopheryl Acetate, Water, Yellow 5.

Store between 15-30C (59-86F)  
Avoid freezing and excessive heat above 40C (104F)

Call toll free 1-877-274-8358

## Packaging

218.8mm×25mm×167mm



Note: This mock-up is for **reference only** and is not the actual artwork properly. **Please refer to the actual artwork properly.** Proper graphic alignment is required.

alcohol spray

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-541
Route of Administration	TOPICAL		

Ingredient Name	Basis of Strength	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients				
Ingredient Name			Strength	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)				
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)				
FD&C RED NO. 4 (UNII: X3W0AM1JLX)				
WATER (UNII: 059QF0KO0R)				
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
GLYCERIN (UNII: PDC6A3C0OX)				
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-541-01	3 in 1 BOX	01/04/2026	
1		50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M003	01/04/2026	

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

Establishment			
Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-541)

Revised: 1/2026

Shenzhen Lantern Science Co.,Ltd.