

SMART CARE HAND SANITIZER 3PK - FLORAL FRESH (LAVENDER/WILD ROSE/RAIN)- alcohol spray
Shenzhen Lantern Science Co.,Ltd.

Smart Care Hand Sanitizer 3pk - Floral Fresh (Lavender/Wild rose/Rain)

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.

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 33, Tocopheryl Acetate, Water, Yellow 5.

other Information

Store between 15-30C (59-86F)

Avoid freezing and excessive heat above 40C (104F)

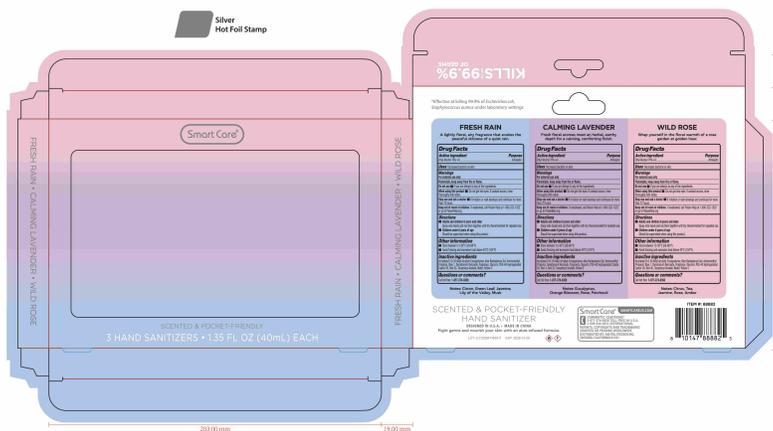
Spray 3-4 times each time.

Used in the skin area of the hands.

packing

Packaging

Brand DC	Quantity: 3pk	Designer: KCal	Notes
Sub Brand: Subtoprim	Category: Packaging	Draw: 11-23	
Item Name: Hand Sanitizer	Item No: 00802	Substrate:	
Description: Floral Fresh	Version: v2	Finish:	



KILLS 99.9% OF GERMS



NOTE: It is for reference only and a reminder that the packaging has feet.

SMART CARE HAND SANITIZER 3PK - FLORAL FRESH (LAVENDER/WILD ROSE/RAIN)

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-537
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
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0K00R)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C00X)	
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:54860-537-01	3 in 1 BOX	12/03/2025	
1		40 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	12/03/2025	

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

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-537)

Revised: 12/2025

Shenzhen Lantern Science Co.,Ltd.