

**SMART CARE LUXURY HAND SANITIZER 1.35OZ - CITRUS SUEDE-
alcohol spray
Shenzhen Lantern Science Co.,Ltd.**

Smart Care Luxury Hand Sanitizer Spray 1.35oz - Citrus Suede

Drug Facts

Active Ingredient

Active ingredient Purpose

Ethyl Alcohol 70% v/v Antiseptic

Uses

Decreases bacteria on skin

Warning

For external use only

Flammable, keep away from fire or flame.

When using this product

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

Do not use

If you are allergic to any of the ingredients

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

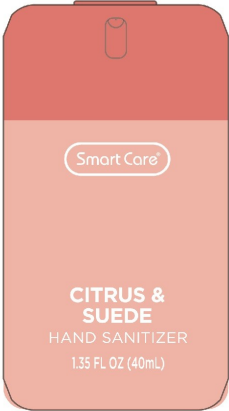
Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Gel, Aminomethyl Propanol,Buiue 1, Denatonium Benzoate, Fragrance, Glycerin, Peg-40 Hydrogenated Castor Oil, Red 4, Red 33, Tocopheryl Acetate, Water.

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

packing




Front View




Back View

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


White Sticker Label Booklet (Printing)




BLACK PANTONE 2030 C




Back Sticker (cover)



Drug Facts (Continued)



Drug Facts (Continued)



Drug Facts (Continued)

SMART CARE LUXURY HAND SANITIZER 1.35OZ - CITRUS SUEDE			
alcohol spray			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-464
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)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-464-01	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 Science Co.,Ltd. (421222423)

Establishment			
Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-464)

Revised: 4/2025

Shenzhen Lantern Science Co.,Ltd.