

SMART CARE HAND SANITIZER PINE FOREST- alcohol spray
Shenzhen Lantern Science Co.,Ltd.

Smart Care Hand Sanitizer Spray 1.35oz (40mL) - Pine Forest

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, Blue 1, Denatonium Benzoate, Fragrance, Glycerin, Peg-40 Hydrogenated Castor Oil, Red 33, Tocopheryl Acetate, Water, Yellow 5.

other Information

Store between 15-30°C (59-86°F)

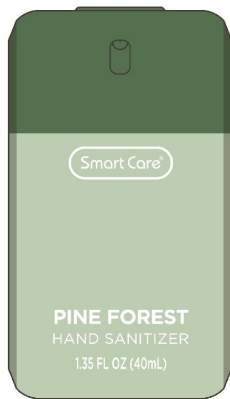
Avoid freezing and excessive heat above 40°C (104°F)

Spray 3-4 times each time.

Used in the skin area of the hands.

packing

packing



Front View



Back View

White Sticker Label Booklet (Printing)

BLACK 2048 C
44mm
63.76mm

Back Sticker (Cover)

Back Sticker Page 2 (Inside cover)

Back Sticker Page 3 (Inside cover)

Back Sticker Page 4 (Inside cover)

SMART CARE HAND SANITIZER PINE FOREST

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-470
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
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
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 BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-470-01	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(54860-470)

Revised: 4/2025

Shenzhen Lantern Science Co.,Ltd.