

SMART CARE HAND SANITIZER-SQUISHMALLOWS MAUI- alcohol spray
Shenzhen Lantern Science Co.,Ltd.

Smart Care Hand Sanitizer-Squishmallows Maui

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, Denatonium Benzoate, Fragrance, Glycerin, PEG-40 Hydrogenated Castor Oil, Red 4, Tocopheryl Acetate, Water, Yellow 5.

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

	Designer: NW-02.ai	Notes:
19	Date: 4.18.25	
	Substrate:	
	Finish:	


shell:  Color of Hand Sanitizer Liquid: 

PANTONE 100C PANTONE 120 C








Front View




Back View

White Sticker Label Booklet (Printing)




Back Sticker (Cover)




Back Sticker Page 2 (Inside cover)



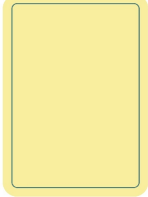
Back Sticker Page 3 (Inside cover)



Back Sticker Page 4 (Inside cover)



Back Sticker Page 5 (Inside cover)



Back Adhesive Side (This color will be seen from front)

SMART CARE HAND SANITIZER-SQUISHMALLOWS MAUI

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-483
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
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
WATER (UNII: 059QF0KO0R)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-483-01	50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/23/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/23/2025	

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

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-483)

Revised: 6/2025

Shenzhen Lantern Science Co.,Ltd.