

SMART CARE HAND SANITIZER - BARBIE- 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 33, Tocopheryl Acetate, Water.

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



White Sticker Label Booklet (Printing)

BLACK

PANTONE 805C

Peony Petals Scent

Peel ▶

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ITEM #: 88439

10147 88439 1

MADE IN CHINA

LOT: JLT250032851902

EXP: 2025-06-25

Drug Facts (Continued)

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*Effective at killing 99.9% of *Escherichia coli*, *Staphylococcus aureus* under laboratory settings

Drug Facts (Continued)

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Drug Facts (Continued)

Questions or comments?
Call toll free 1-877-274-8388

Smart Care SMARTCAREUS.COM

QUESTIONS OR COMMENTS?
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1-800-834-8811 INTERNATIONAL
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Back Sticker (cover)

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

SMART CARE HAND SANITIZER - BARBIE

alcohol spray

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:54860-485
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)				
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-485-01	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(54860-485)