SMART CARE HAND SANITIZER - BARBIE- alcohol spray
Shenzhen Lantern Scicence Co.,Ltd.

Smart Care Hand Sanitizer-Squishmallows Maui

Drug Facts

Active Ingredient

Active ingredient Purpose

Ethyl Alcohol 70% v/v Antiseptic

Uses

Decreases bacieria on skin

Warning

For external use only

Flammable, keep away from fire or flame.

When using this product

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

Do not use

lf you are allergic to any of the ingredients

Stop use and ask a doctor

lf irritation or rash develops and continues for more than72 hours.

keep out of reach of children

lf 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 unil 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.

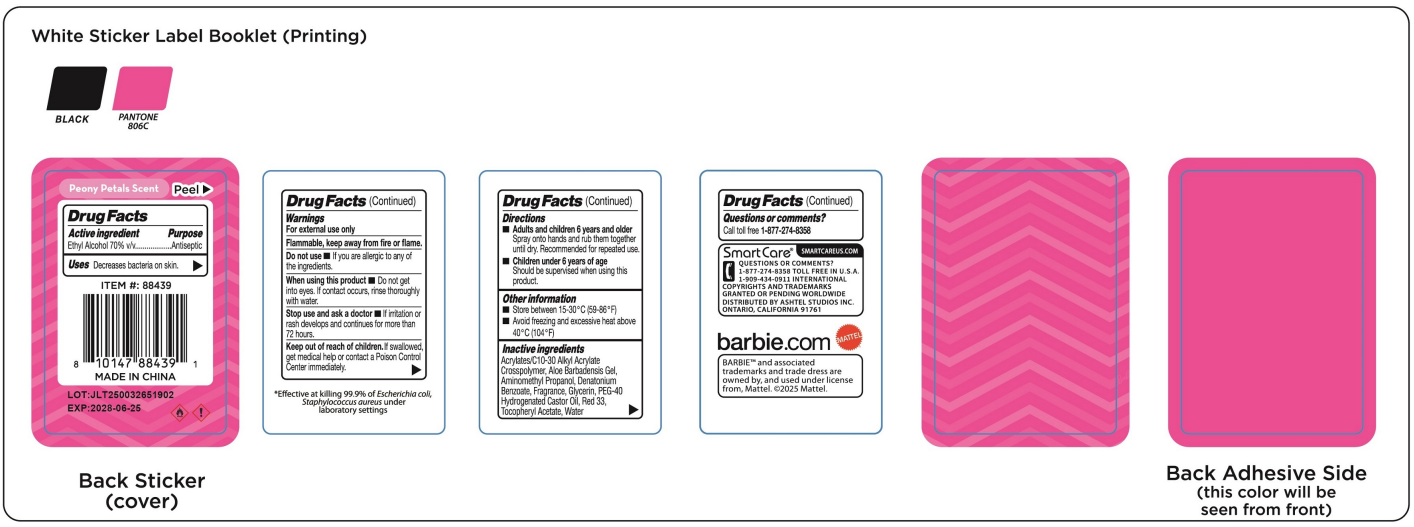
other Information

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



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 Scicence Co.,Ltd. (421222423)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-485)

Revised: 6/2025 Shenzhen Lantern Scicence Co.,Ltd.