

SMART CARE HAND SANITIZER MONSTER JAM- alcohol spray
Shenzhen Lantern Science Co.,Ltd.

Smart Care Hand Sanitizer Monster Jam

Drug Facts

Active Ingredient

Active ingredient Purpose

Ethyl Alcohol 70% v/v Antiseptic

Uses

Decreases bacteria on skin

Warning

For external use only

Do not use

If you are allergic to any of the ingredients

When using this product

Do not get into eyes. If contact occurs, rinse thoroughly with water. Do not use in or near eyes. In case of eye contact, flush eyes thoroughly with water. Discontinue if skin becomes irritated and ask for a doctor.

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, Aminomethy Propanol, Denatonium Benzoate, Fragrance, Glycerin, Tocopheryl Acetate, Water.

other Information

Storebetween15-30C(59-86F)
Avoid freezing and excessive heat above 40C (104F)

Questions or comments?

Call toll free 1-877-274-8358

packaging

Packagin



SMART CARE HAND SANITIZER MONSTER JAM
alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-551
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
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)				
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-551-01	28 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/05/2026	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M003	01/05/2026	

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

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-551)