LICENSED HAND SANITIZER SPIDER MAN 01- alcohol spray
Shenzhen Lantern Scicence Co.,Ltd.

88113 Licensed Hand Sanitizer Spray Spider man

Drug Facts

Active Ingredient

Active ingredient Purpose

Ethyl Alcohol 70% v/v Antiseptic

Uses

Decreases bacieria on skin

Warning

For external use only

Do not use

lf you are allergic to any of the ingredients

When using this product

Do not get into eyes. lf 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

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

Water, Fragrance,Glycerin, ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER, Aminomethyl Propanol, Tocopheryl Acetate, Aloe Barbadensis Gel, Denatonium Benzoate.

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

146x127x31mm

Inside Background

LICENSED HAND SANITIZER SPIDER MAN 01

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54860-435
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)	
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-435-01	3 in 1 BOX	02/14/2025	
1		28 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	02/14/2025	

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

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-435)

Revised: 2/2025 Shenzhen Lantern Scicence Co.,Ltd.