

**LICENSED HAND SANITIZER HOT WHEELS 01- alcohol spray
Shenzhen Lantern Science Co.,Ltd.**

88111 Licensed Hand Sanitizer Spray Hot Wheels

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

Water, Fragrance, Glycerin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Tocopheryl Acetate, Aloe Barbadensis Gel, Denatonium Benzoate.

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

Packaging

146x127x31mm

Inside Background



LICENSED HAND SANITIZER HOT WHEELS 01

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-433
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)	
WATER (UNII: 059QF0K00R)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-433-01	3 in 1 BOX	02/12/2025	
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	02/12/2025	

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

Establishment			
Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-433)

Revised: 2/2025

Shenzhen Lantern Science Co.,Ltd.