SMART CARE LICENSED HAND SANITIZER HOT WHEELS PURPLE- alcohol spray
Shenzhen Lantern Scicence Co.,Ltd.

Smart Care Licensed Hand Sanitizer Hot Wheels Purple

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

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Gel, Aminomethy Propanol, Denatonium Benzoate, Fragrance, Glycerin, PEG-40 Hydrogenated Castor Oil, Tocopheryl Acetate, Water.

other Information

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

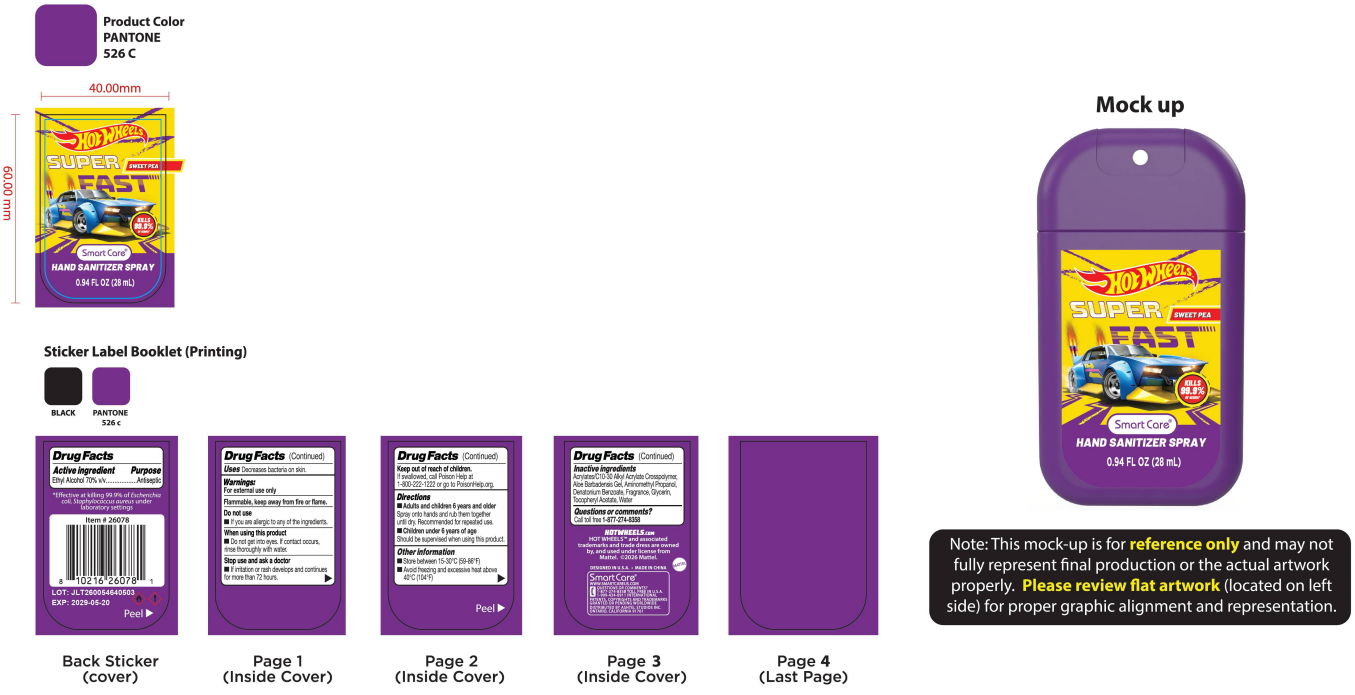
Questions or comments?

Call toll free 1-877-274-8358

packaging

Packagin

Brand: SC	**Quantity:** 3pk	**Designer:** KC.ai	**Notes:**
Sub brand: HotWheels	**Category:** products	**Date:** 3.5.26	
Item Name: Hand Sanitizer	**Item No:** 26078	**Substrate:**	
Description: 0.94oz	**Version:** v1	**Finish:**	



SMART CARE LICENSED HAND SANITIZER HOT WHEELS PURPLE

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54860-567
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
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)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-567-01	28 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/08/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	04/08/2026	

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

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen Lantern Science Co.,Ltd.		421222423	manufacture(54860-567)

Revised: 4/2026 Shenzhen Lantern Scicence Co.,Ltd.