

HOT WHEELS HAND SANITIZER- ethyl alcohol gel
Ashtel Studios, Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Hot Wheels HAND SANITIZER

Drug Facts

Active Ingredients

Ethyl Alcohol 62% v/v

Purpose

Antiseptic

Use:

To help reduce bacteria and germs on the skin.

WARNINGS:

For external use only: hands.

Flammable. Keep away from fire or flame.

When using this product:

• Keep out of eyes. • In case of contact with eyes, flush thoroughly with water. • Avoid contact with broken skin. • Do not inhale or ingest.

• **Keep out of reach of children.**

• In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions:

Put a thumbnail size amount in your palm and rub your hands together briskly until dry. Children under 6 years of age should be supervised when using this product. Not recommended for infants.

Other Information:

• Do not store above 100° F (38°C)

• May discolor some fabrics.

• Harmful to wood finishes & plastics.

Inactive Ingredients:

Water (Aqua), Glycerin, Propylene Glycol, Carbomer, Triethanolamine, Aloe Barbadensis Leaf Extract, Fragrance, Tocopheryl Acetate (Vitamin E), FD&C Blue No. 1

QUESTIONS OR COMMENTS?

1-877-274-8358 Toll Free in USA

1-909-434-0911 International

KILLS 99% OF MOST COMMON GERMS

HOTWHEELS.COM

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ASHTEL STUDIOS INC.

ONTARIO, CA 91761

SMARTCAREUS.COM

Packaging



HOT WHEELS HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:70108-061

Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
	Ingredient Name	Basis of Strength	Strength	
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL	
Inactive Ingredients				
	Ingredient Name	Strength		
	WATER (UNII: 059QF0K00R)			
	GLYCERIN (UNII: PDC6A3C0OX)			
	PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
	CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
	TROLAMINE (UNII: 9O3K93S3TK)			
	ALOE VERA LEAF (UNII: ZY81Z83H0X)			
	.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)			
	FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70108-061-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/23/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	06/23/2020		

Labeler - Ashtel Studios, Inc (148689180)