

HOT WHEELS HAND SANITIZER- ethyl alcohol gel
Yozzi Co.,ltd

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 Barbadosensis 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

HOTWHEELS and associated trademarks and trade dress are owned by, and used under licence from Mattel.

© 2020 Mattel. All Rights Reserved.

DESIGNED IN U.S.A.

MADE IN CHINA

PATENTS, COPYRIGHTS AND TRADEMARKS GRANTED OR PENDING WORLDWIDE

DISTRIBUTED BY

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:72285-009

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: 059QF0KO0R)			
	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: HBR47K3TBD)			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72285-009-01	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 - Yozzi Co.,ltd (560426393)

Establishment

Name	Address	ID/FEI	Business Operations
Yozzi Co.,ltd		560426393	manufacture(72285-009)

Revised: 6/2020

Yozzi Co.,ltd