HOT WHEELS HAND SANITIZER- benzalkonium chloride liquid
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 Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antiseptic

Use To help reduce bacteria and germs on the skin.

WARNING Flammable. Keep away from fire or flame. For external use only • Stop use and ask a doctor if irritation or redness develops and persists.

• **Keep out of reach of children.** • In case of accidental digestion, seek professional assistance or contact a Poison Control Center immediately.

Directions • Place enough product in palm to cover hands and rub hands together briskly until dry.

• Children under 6, use only under adult supervision.

• Not recommended for infants.

Other Information • Do not store above 100°F (38° C). • May discolor some fabrics. • Harmful to wood finishes and plastics.

Inactive Ingredients • Aqua (Water), Hydroxyethylcellulose, Phenoxyethanol, Disodium EDTA, Polysorbate 20, Parfum (Fragrance).

HOTWHEELS.COM

QUESTIONS OR COMMENTS ?

1-877-274-8358 Toll Free in USA + 1-909-434-0911 International COPYRIGHTS AND TRADEMARKS GRANTED OR PENDING WORLDWIDE

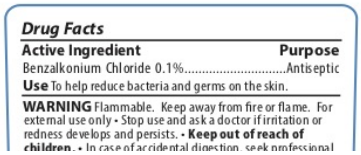
DISTRIBUTED BY ASHTEL STUDIOS INC. ONTARIO, CALIFORNIA 91761

Smart Care®

Packaging

FRONT PANEL

BACK PANEL



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HOT WHEELS HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70108-025
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
HYDROXYETHYL CELLULOSE (4000 MPA.S AT 1%) (UNII: ZYD53NBL45)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70108-025-01	53 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/01/2019	

Labeler - Ashtel Studios, Inc (148689180)

Revised: 10/2019 Ashtel Studios, Inc