

MINIONS 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.

MINIONS 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).

DESPICABLE ME

MINION MADE

KILLS 99% OF GERMS

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QUESTIONS or COMMENTS?

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DISTRIBUTED BY ASHTEL STUDIOS INC. ONTARIO, CALIFORNIA 91761

Smart Care®

SMARTCAREUS.COM

Packaging

FRONT PANEL



BACK PANEL

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benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70 108-023
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 MP.A.S AT 1%) (UNII: ZYD53NBL45)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETATE DISODIUM ANHYDRO US (UNII: 8NLQ36F6MM)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70 108-023-01	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	

