INSTANT HAND SANITIZER- benzalkonium chloride gel Fun Zone, 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.

Instant Hand Sanitizer

Active ingredient

Benzalkonium Chloride 0.095%

Purpose

Antiseptic

Use

To help reduce bacteria on hands.

Warnings

For external use only.

When using this product

- Avoid contact with eyes and lips.
- If contact occurs, rinse with water.

Stop using this product and consult a doctor If

irritation or redness develops.

Keep out of reach of children

If swallowed, get medical help or contact a poison control center immediately.

Directions

- Rub dime sized amount between hands until dry.
- For use by children under 6 years, adult supervision is required.

Inactive ingredients

Water, Polyquaternium-37, Glycerin, Propylene Glycol, Phenoxyethanol, PEG-40 Hydrogenated Castor Oil, Fragrance, Triethanolamine.









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Inactive ingredients Water, Polyquaternium-37, Glycerin, Propylene Glycol, Phenoxyethanol, PEG-40 Hydrogenated Castor Oil, Fragrance, Triethanolamine.

Manufactured for Fun Zone Inc Los Angeles, CA 90067 Made in Rugao City, China

LOT:09/2020 EXP:09/2022

INSTANT HAND SANITIZER

benzalkonium chloride gel

Product Information

HUMAN OTC DRUG NDC:69583-029 Product Type Item Code (Source)

TOPICAL **Route of Administration**

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM -BENZALKONIUM 0.95 mg UNII:7N6JUD5X6Y) CHLORIDE in $1\,\text{mL}$

Inactive Ingredients	
Ingredient Name	Strength
PHENO XYETHANO L (UNII: HIE492ZZ3T)	
POLYQUATERNIUM-37 (10000 MPA.S) (UNII: 41QWS48DFN)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6 A3C0 OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

l	Packaging					
	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1 NDC:69583-029- 01	29 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/04/2020			
П						

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333E	09/04/2020				

Labeler - Fun Zone, Inc. (803572929)

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