SPIDERMAN WATERMELON HAND SANITIZER- benzalkonium chloride gel Shenzhen Lantern Science 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.

Active Ingredient

Benzalkonium Chloride 0.1%

□ **Purpose:** □ Antibacterial

 \Box Use

To decrease bacteria on the skin that could cause disease.

Keep out of reach of children.

□ Warnings

- For external use only-hands.
- Keep out of eyes. Avoid contact with broken skin.
- **Stop use and ask a doctor if** irritation or redness develops.
- **Do not inhale or ingest.** If swallowed, get medical help or contact a poison control center right away.

Directions

- Rub a dime sized drop into hands.
- For children under 6 use under adult supervision.

Inactive Ingredients

water (aqua/eau), glycerin, coceth-7, PPG-1-PEG-9 lauryl glycol ether, carbomer, hydrogenated castor oil, fragrance (parfum).

> CMYK PROCESS CLEAR ADHESIVE STOCK

CONFIRM WEIGHTS + FLAVORS + WARNING

May Contain

Red 40 (CI 16035), Red 33 (CI 17200), Blue 1 (CI 42090), Yellow 5 (CI 19140).

Hand Sanitizer





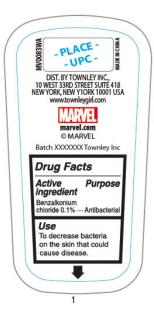
















SPIDERMAN WATERMELON HAND SANITIZER

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-072
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	0.059 g in 59 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
GLYCERIN (UNII: PDC6 A3C0 OX)			
COCETH-7 CARBOXYLIC ACID (UNII: 35KO064932)			
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)			
PEG-40 CASTOR OIL (UNII: 4ERD2076EF)			
PPG-1-PEG-9 LAURYL GLYCOL ETHER (UNII: 5R8J43K25L)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			

ı	Packagin	ıg			
ı	# Item	Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:548	60-072-01 59 i	mL in 1 BOTTLE; Type 0: Not a Combination Product	0 1/18/20 17	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	0 1/18/20 17		

Labeler - Shenzhen Lantern Science Co., Ltd. (421222423)

Establishment				
Name	Address	ID/FEI	Business Operations	
Shenzhen Lantern Science Co., Ltd.		421222423	manufacture (54860-072)	

Revised: 1/2017 Shenzhen Lantern Science Co., Ltd.