

**MARVELS ULTIMATE SPIDER MAN STRAWBERRY SCENTED 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.

PURPOSE

Antibacterial

To decrease bacteria on skin that could cause disease

ACTIVE INGREDIENT

BENZALKONIUM CHLORIDE.....0.1%

DOSAGE AND ADMINISTRATION

Rub a dime sized drop into hands

For children under 6 use under adult supervision

WARNINGS

For external use only - hands

Keep out of eyes.

Avoid contact with broken skin.

Do not inhale or ingest

If swallowed get medical help or contact a poison control center right away

KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children

For children under 6, use under adult supervision

INDICATIONS AND USAGE

To decrease bacteria on skin that could cause disease.

INACTIVE INGREDIENTS

WATER (AQUA/EAU)

GLYCERIN

COCETH-7

PPG-1-PEG-9 LAURYL GLYCOL ETHER

CARBOMER

PEG-40 HYDROGENATED CASTOR OIL

FRAGRANCE (PARFUM)

SODIUM HYDROXIDE

PACKAGE LABEL

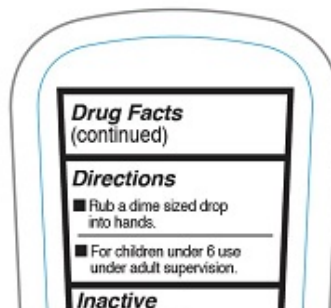
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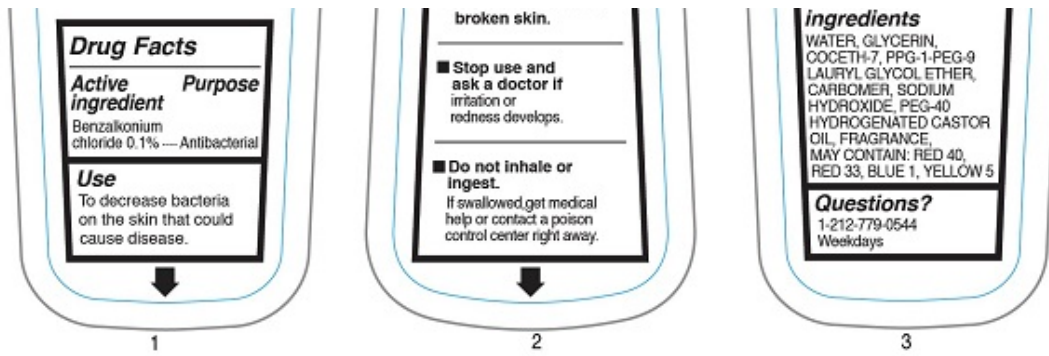
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**CONFIRM WEIGHTS +
 FLAVORS + WARNING
 White Base X2**

Townley
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 New York, NY 10016
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MARVELS ULTIMATE SPIDER MAN STRAWBERRY SCENTED HAND SANITIZER

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54860-021
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 100 mg

Inactive Ingredients	
Ingredient Name	Strength
CO CETH-7 (UNII: 58 Y261JLH5)	
CARBOMER 1342 (UNII: 809 Y72KV36)	
GLYCERIN (UNII: PDC6 A3C0OX)	
PPG-1-PEG-9 LAURYL GLYCOL ETHER (UNII: 5R8J43K25L)	
SODIUM HYDRO XIDE (UNII: 55X04QC32I)	
CASTOR OIL (UNII: D5340 Y2I9G)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color	red (RED 40) , red (RED 33) , blue (BLUE 1) , yellow (YELLOW 5)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54860-021-01	60 mg in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/14/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/14/2016	

Labeler - SHENZHEN LANTERN SCIENCE CO., LTD (421222423)

Establishment

Name	Address	ID/FEI	Business Operations
SHENZHEN LANTERN SCIENCE CO., LTD		421222423	manufacture(54860-021)

Revised: 3/2016

SHENZHEN LANTERN SCIENCE CO., LTD