AVENGERS MARVEL HAND SANITIZER- benzalkonium chloride gel
Townley, 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.

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose: Antibacterial

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).

May Contain

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

Avengers Marvel Hand Sanitizer

AVENGERS MARVEL HAND SANITIZER

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58737-162
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 g in 59 g

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
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)	

Product Characteristics			
Color	red (red 40) , red (Red 33) , blue (Blue 1) , yellow (Yellow 5)	**Score**	
Shape		**Size**	
Flavor	BLUEBERRY, CHERRY, LEMON, BERRY, GRAPE	**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:58737-162-01	59 g in 1 BOTTLE; Type 0: Not a Combination Product	08/10/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/10/2016	

Labeler - Townley, Inc. (016956158)

Establishment

Name	Address	ID/FEI	Business Operations
Townley, Inc.		016956158	manufacture(58737-162)