

**BUBBLEGUM SCENTED 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.*

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***Active Ingredient***

Benzalkonium Chloride 0.1%

***Purpose***

Antibacterial

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***

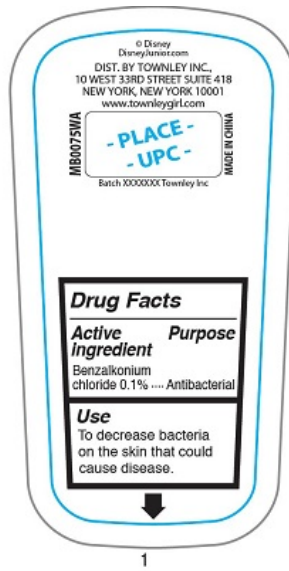
- Rub a dime sized drop into hands.
- For children under 6 use under adult supervision.

***Inactive Ingredients***

water, glycerin, coceth-7, PPG-1-PEG-9 lauryl glycol ether, carbomer, sodium hydroxide, PEG-40 hydrogenated castor oil, fragrance

***Use***

To decrease bacteria on the skin that could cause disease.



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## BUBBLEGUM SCENTED HAND SANITIZER

benzalkonium chloride gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58737-131
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>COCETH-7</b> (UNII: 58Y261JLH5)	
<b>PPG-1-PEG-9 LAURYL GLYCOL ETHER</b> (UNII: 5R8J43K25L)	
<b>CARBOMER COPOLYMER TYPE A</b> (UNII: 71DD5V995L)	
<b>POLYOXYL 40 HYDROGENATED CASTOR OIL</b> (UNII: 7YC686GQ8F)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58737-131-01	59 g in 1 BOTTLE; Type 0: Not a Combination Product	05/16/2016	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/16/2016	

**Labeler** - Townley, Inc. (016956158)

## Establishment

Name	Address	ID/FEI	Business Operations
Foshan Jinxiong Technology Co., Ltd		544328419	manufacture(58737-131)

Revised: 5/2016

Townley, Inc.