# BLUEBERRY 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.

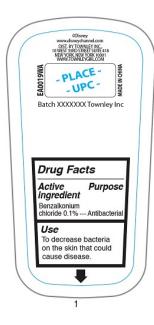
## **Inactive Ingredients**

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

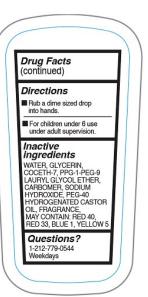
#### Use

To decrease bacteria on the skin that could cause disease.









## **BLUEBERRY SCENTED HAND SANITIZER**

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58737-129
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
<b>BENZALKO NIUM CHLO RIDE</b> (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g		

Inactive Ingredients	
Ingredient Name	Strength
SO DIUM HYDRO XIDE (UNII: 55X0 4QC32I)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CO CETH-7 (UNII: 58 Y26 1JLH5)	
PPG-1-PEG-9 LAURYL GLYCOL ETHER (UNII: 5R8J43K25L)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	

F	Packaging			
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>

1 NDC:58737-129-01	59 g in 1 BOTTLE; Type 0: Not a Combination Product	05/16/2016		
Marketing Information				
Marketing Categor	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not fina	al part333A	05/16/2016		

## Labeler - Townley, Inc. (016956158)

Establishment				
Name	Address	ID/FEI	Business Operations	
Foshan Jinxiong Technology Co., Ltd		544328419	manufacture(58737-129)	

Revised: 5/2016 Townley, Inc.