MY LITTLE PONY COTTON CANDY SCENTED- 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

Directions

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

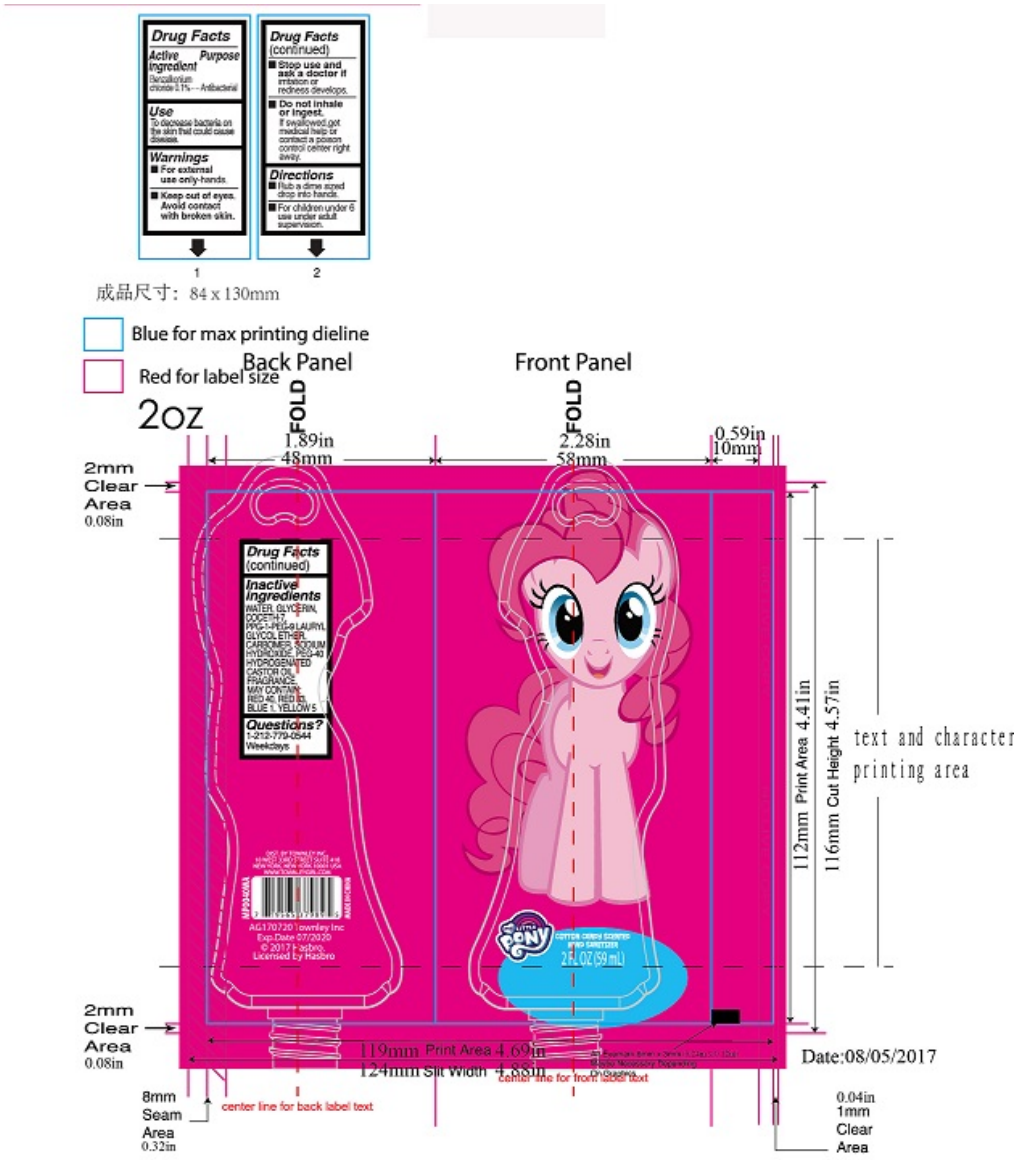
Warnings

- **For external use only -** hands.
- **Keep out of eyes. Avoid contact with broken skin.**
- **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.

keep out of reach of children

Inactive ingredients

water, glycerin, coceth-7, PPG-1-PEG-9 lauryl glycol ether, carbomer, sodium hydroxide, PEG-40 hydrogenated castor oil, fragrance, may contain: red 40, red 33, blue 1, yellow 5.



MY LITTLE PONY COTTON CANDY SCENTED

benzalkonium chloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58737-219
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	

POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCETH-7 (UNII: 58Y261JLH5)	
PPG-1-PEG-9 LAURYL GLYCOL ETHER (UNII: 5R8J43K25L)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	COTTON CANDY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58737-219-01	59 g in 1 BOTTLE; Type 0: Not a Combination Product	08/22/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/22/2017	

Labeler - Townley Inc. (016956158)

Establishment

Name	Address	ID/FEI	Business Operations
Foshan Jinxiong Technology Co ., Ltd.		544328419	manufacture(58737-219)

Revised: 8/2017 Townley Inc.