PERSONAL CARE FOAMING HAND SANITIZER- benzalkonium chloride liquid
Delta Brands & Products LLC

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.

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antibacterial

Uses

for washing hands to decrease bacteria on the skin

Warnings

For external use only

When using this product

■ avoid contact with the eyes. In case of eye contact, flush with water.

Stop use and ask a doctor if

■ irritation or rash develops

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

■ pump into hands ■ lather vigorously for at least 15 seconds ■ rinse and dry thoroughly.

Inactive ingredients

water (aqua), laureth-9, disodium cocoamphodiacetate, polyquaternium-10, citric acid, fragrance, glycerin, methylchloroisothiazolinone, methylisothiazolinone, FD&C blue no. 1, FD&C yellow no. 5

Package Label

PERSONAL CARE FOAMING HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72133-150
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 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
EDETATE SODIUM TETRAHYDRATE (UNII: L13NHD21X6)	
WATER (UNII: 059QF0KO0R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLIDOCANOL (UNII: 0AWH8BFG9A)	

POLYQUATERNIUM-10 (1000 MPA.S AT 2%) (UNII: GMR4PEN8PK)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72133-150-75	221.8 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/02/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/02/2018	

Labeler - Delta Brands & Products LLC (080999173)

Revised: 4/2018 Delta Brands & Products LLC