LUCKY ANTIBACTERIAL FOAMING HANDWASH- benzalkonium chloride liquid
Delta Brands, 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.

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

to decrease bacteria on the skin

Warnings

For external use only

■ avoid eye contact. In case of eye contact, flush with water. If irritation occurs discontinue use of product

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 and rinse

Inactive ingredients

water, cocamidopropyl betaine, decyl glucoside, glycerin, fragrance, citric acid, polyquaternium-7, xanthan gum, methylchloroisothiazolinone, methylisothiazolinone, polysorbate-20, tetrasodium EDTA, aloe barbadensis leaf juice, FDC blue no. 1, DC red no. 33

Package Label

LUCKY ANTIBACTERIAL FOAMING HANDWASH

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:20276-143
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	

DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
XANTHAN GUM (UNII: TTV12P4NEE)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:20276-143-75	221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/2017	

Labeler - Delta Brands, Inc (102672008)

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